

**The Osmoadaptive Response of the Wine Yeast *Saccharomyces cerevisiae* K1-V1116
during Icewine Fermentation**

Stephanie J. Martin, B.Sc. Hon.

Submitted to the Centre for Biotechnology in partial fulfillment of the requirements for
the degree of Doctor of Philosophy

JAMES A GIBSON LIBRARY
BROCK UNIVERSITY
ST. CATHARINES ON

August 2008

Brock University

St. Catharines, Ontario

©Stephanie J. Martin, 2008

ABSTRACT

The adapted metabolic response of commercial wine yeast under prolonged exposure to concentrated solutes present in Icewine juice is not fully understood. Presently, there is no information regarding the transcriptomic changes in gene expression associated with the adaptive stress response of wine yeast during Icewine fermentation compared to table wine fermentation. To understand how and why wine yeast respond differently at the genomic level and ultimately at the metabolic level during Icewine fermentation, the focus of this project was to identify and compare these differences in the wine yeast *Saccharomyces cerevisiae* K1-V1116 using cDNA microarray technology during the first five days of fermentation. Significant differences in yeast gene expression patterns between fermentation conditions were correlated to differences in nutrient utilization and metabolite production. Sugar consumption, nitrogen usage and metabolite levels were measured using enzyme assays and HPLC. Also, a small subset of differentially expressed genes was verified using Northern analysis. The high osmotic stress experienced by wine yeast throughout Icewine fermentation elicited changes in cell growth and metabolism correlating to several fermentation difficulties, including reduced biomass accumulation and fermentation rate. Genes associated with carbohydrate and nitrogen transport and metabolism were expressed at lower levels in Icewine juice fermenting cells compared to dilute juice fermenting cells. Osmotic stress, not nutrient availability during Icewine fermentation appears to impede sugar and nitrogen utilization. Previous studies have established that glycerol and acetic acid production are increased in yeast during Icewine fermentation. A gene encoding for a glycerol/H⁺ symporter (*STL1*) was found to be highly expressed up to 25-fold in the

Icewine juice condition using microarray and Northern analysis. Active glycerol transport by yeast under hyperosmotic conditions to increase cytosolic glycerol concentration may contribute to reduced cell growth observed in the Icewine juice condition. Additionally, genes encoding for two acetyl CoA synthetase isoforms (*ACS1* and *ACS2*) were found to be highly expressed, 19- and 11-fold respectively, in dilute juice fermenting cells relative to the Icewine juice condition. Therefore, decreased conversion of acetate to acetyl-CoA may contribute to increased acetic acid production during Icewine fermentation. These results further help to explain the response of wine yeast as they adapt to Icewine juice fermentation.

ACKNOWLEDGEMENTS

I would like to personally thank my supervisor Dr. Debra Inglis for providing me the opportunity to be a part of a wonderful lab group. Your guidance and encouragement allowed me to develop and more importantly, have confidence in, my research skills. I truly appreciate all of your help over the years and I am thrilled that I have been one of your students.

I also want to thank Dr. Gary Pigeau, a very talented researcher whose enviable technical skills and academic expertise always set me in the right direction.

To everyone in the IH210 lab group- past and present members (Ai Lin, Shiri, Qing, Fei, Heidi, Lisa, Eric, Wendy and Matilda)- you are a fantastic collection of exceptionally talented people and all of whom are extremely funny, sweet and best of all, supportive. Each of you have influenced my life in so many amazing ways and for that, I thank you all.

I also want to extend my deepest appreciation and thanks for all the faculty and staff of CCOVI, Department of Biology and Department of Chemistry I had the privilege of working with throughout my graduate studies.

Last, but definitely not least, I would like to thank my mother and father for their continuing love, support, guidance and patience. Thank you both for always giving me the drive to pursue my dreams.

TABLE OF CONTENTS

| | |
|---|------------|
| ABSTRACT | i |
| ACKNOWLEDGEMENTS | iii |
| TABLE OF CONTENTS | iv |
| LIST OF TABLES | vi |
| LIST OF FIGURES | vii |
| 1. INTRODUCTION..... | 8 |
| 1.1. INTRODUCTION TO THE PROBLEM..... | 8 |
| 1.2. OBJECTIVES | 9 |
| 1.3. EXPERIMENTAL DESIGN | 10 |
| 2. LITERATURE REVIEW | 11 |
| 2.1. INTRODUCTION TO ICEWINE: PRODUCTION AND CHALLENGES | 11 |
| 2.2. RESPONDING TO HYPEROSMOTIC STRESS AND OSMOADAPTATION IN <i>S. CEREVISIAE</i> | 14 |
| 2.2.1. Hyperosmotic stress: cellular effects, consequences and responses | 14 |
| 2.2.2. Signaling pathways involved in the hyperosmotic stress response..... | 15 |
| 2.2.2.1. <i>The HOG response</i> | 15 |
| 2.2.2.2. <i>The general stress response</i> | 21 |
| 2.2.2.3. <i>Immediate response to hyperosmotic stress</i> | 22 |
| 2.2.3. Glycerol metabolism in <i>S. cerevisiae</i> | 23 |
| 2.2.3.1. <i>Glycerol synthesis during hyperosmotic stress</i> | 24 |
| 2.2.3.2. <i>Glycerol retention and uptake</i> | 28 |
| 2.2.4. Acetic acid metabolism in <i>S. cerevisiae</i> | 31 |
| 2.2.5. Pyruvate dehydrogenase bypass | 33 |
| 2.3. GENE EXPRESSION PROFILING IN <i>S. CEREVISIAE</i> DURING FERMENTATION | 35 |
| 2.3.1. Gene expression of laboratory yeast strains under salt stress | 36 |
| 2.3.2. Gene expression profiling in wine yeast | 38 |
| 2.4. YEAST SUGAR AND NITROGEN METABOLISM | 40 |
| 2.4.1. Sugar Utilization | 40 |
| 2.4.2. Nitrogen Utilization..... | 45 |
| 3. MATERIALS AND METHODS | 49 |
| 3.1. YEAST STRAINS..... | 49 |
| 3.2. MEDIA | 49 |
| 3.3. ICEWINE JUICE PREPARATION AND COMPOSITIONAL ANALYSIS..... | 50 |
| 3.4. STARTER CULTURE PREPARATION AND INOCULATION | 52 |
| 3.5. FERMENTATION AND METABOLITE ANALYSIS | 53 |
| 3.6. RNA EXTRACTION | 54 |
| 3.7. MICROARRAY ANALYSIS | 55 |
| 3.7.1. Reverse transcription and hybridization..... | 55 |
| 3.7.2. Image acquisition and data analysis | 56 |
| 3.8. VALIDATION OF MICROARRAY RESULTS USING NORTHERN ANALYSIS | 58 |
| 3.8.1. DNA Probe Design..... | 58 |
| 3.8.2. Northern Analysis | 59 |
| 3.9. MEASURING AMINO ACID CONCENTRATION USING HPLC | 60 |
| 4. RESULTS AND DISCUSSION | 61 |

| | |
|---|------------|
| 4.1. MICROARRAY ANALYSIS OVERVIEW | 61 |
| 4.1.1. Results - Microarray..... | 61 |
| 4.1.2. Discussion - Microarray Overview | 68 |
| 4.2. CARBOHYDRATE METABOLISM..... | 71 |
| 4.2.1. Results - Carbohydrate Metabolism..... | 71 |
| 4.2.1.1. <i>Yeast Cell Growth</i> | 71 |
| 4.2.1.2. <i>Sugar Consumption</i> | 75 |
| 4.2.1.3. <i>Gene expression profiles for carbohydrate metabolism</i> | 77 |
| 4.2.1.3.1. <i>Glycolysis and alcoholic fermentation</i> | 77 |
| 4.2.1.3.2. <i>TCA and glyoxylate cycles</i> | 82 |
| 4.2.2. Discussion - Sugar Metabolism | 84 |
| 4.3. NITROGEN METABOLISM..... | 89 |
| 4.3.1. Results- Nitrogen Metabolism | 89 |
| 4.3.1.1. <i>Amino acid composition and nitrogen utilization</i> | 89 |
| 4.3.1.2. <i>Gene expression profiles for nitrogen transport and metabolism</i> | 97 |
| 4.3.2. Discussion - Nitrogen Metabolism..... | 105 |
| 4.3.2.1. <i>Amino acid content of Icewine juice</i> | 105 |
| 4.3.2.2. <i>General amino acid transport</i> | 106 |
| 4.3.2.3. <i>Specific amino acid uptake and metabolism</i> | 108 |
| 4.4. GLYCEROL METABOLISM..... | 115 |
| 4.4.1. Results- Glycerol Metabolism..... | 115 |
| 4.4.1.1. <i>Glycerol Production</i> | 115 |
| 4.4.1.2. <i>Gene expression profiles (Microarrays)- Glycerol metabolism and transport</i> | 117 |
| 4.4.1.3. <i>Gene expression profiles (Northern)- Glycerol</i> | 120 |
| 4.4.2. Discussion - Glycerol metabolism and transport | 122 |
| 4.5. ACETIC ACID METABOLISM..... | 126 |
| 4.5.1. Results - Acetic Acid Metabolism | 126 |
| 4.5.1.1. <i>Acetic Acid Production</i> | 126 |
| 4.5.1.2. <i>Gene expression profiles for acetic acid and fatty acid metabolism</i> | 128 |
| 4.5.1.3. <i>Gene expression profiles (Northern)- Acetic acid</i> | 136 |
| 4.5.2. Discussion- Acetic acid Metabolism..... | 141 |
| 4.6. ADDITIONAL TRANSCRIPTOME RESPONSES DURING ICEWINE FERMENTATION | 146 |
| 4.6.1. Results - additional transcriptome responses | 146 |
| 4.6.1.1. <i>Stress Responsive Genes</i> | 146 |
| 4.6.1.2. <i>Transport Responses</i> | 151 |
| 4.6.2. Discussion – additional transcriptome responses..... | 153 |
| 5. OVERALL DISCUSSION AND FUTURE DIRECTIONS | 161 |
| 6. CONCLUSIONS | 170 |
| LITERATURE CITED | 172 |
| APPENDIX I | 188 |
| APPENDIX II..... | 315 |

LIST OF TABLES

| | |
|--|-----|
| Table 3.1: Icewine juice parameters..... | 51 |
| Table 3.2: Primer sequences used for Northern analysis probe design..... | 60 |
| Table 4.1.1: Distribution of the differentially expressed genes in <i>S. cerevisiae</i> fermenting Icelandic wine and dilute Icelandic wine..... | 62 |
| Table 4.1.2: Distribution of GO Slim terms for differentially expressed genes in <i>S.</i> <i>cerevisiae</i> fermenting Icelandic wine..... | 64 |
| Table 4.1.3: Genes up- or downregulated more than 10-fold in <i>S. cerevisiae</i> fermenting Icelandic wine..... | 67 |
| Table 4.2.1: Expression profile of the genes involved in carbohydrate metabolism..... | 80 |
| Table 4.3.1: Amino acids present in unfermented Icelandic wine and finished wine..... | 93 |
| Table 4.3.2: Amino acids present in unfermented diluted Icelandic wine and finished wine..... | 94 |
| Table 4.3.3: Amino acid utilization over time by yeast fermenting Icelandic wine..... | 96 |
| Table 4.3.4: Amino acid utilization over time by yeast fermenting diluted Icelandic wine..... | 96 |
| Table 4.3.5: Changes in the expression profile of genes involved in amino acid metabolism..... | 99 |
| Table 4.4.1: Glycerol produced by <i>S. cerevisiae</i> following the consumption of 187 g L ⁻¹ of sugar..... | 116 |
| Table 4.4.2: Expression profile of the genes involved in glycerol metabolism..... | 119 |
| Table 4.5.1: Metabolites produced by <i>S. cerevisiae</i> following the consumption of 187 g L ⁻¹ of sugar..... | 127 |
| Table 4.5.2: Expression profile of the genes involved in acetic acid metabolism..... | 129 |
| Table 4.5.3: Expression profile of the genes involved in lipid metabolism..... | 132 |
| Table 4.6.1: Expression profile of the genes involved in responding to stress..... | 147 |
| Table 4.6.2: Expression profile of the genes involved in transport..... | 152 |

LIST OF FIGURES

| | |
|--|-----|
| Figure 2.1: The HOG pathway in <i>S. cerevisiae</i> .. | 17 |
| Figure 2.2: Glycerol and acetic acid metabolic pathways in <i>S. cerevisiae</i> .. | 26 |
| Figure 2.3: Pathways associated with carbohydrate metabolism in <i>S. cerevisiae</i> .. | 41 |
| Figure 4.1.1: The major Gene Ontology (GO) Slim term categories containing genes differentially expressed in <i>S. cerevisiae</i> fermenting Icewine juice .. | 65 |
| Figure 4.2.1: Yeast cell morphology during fermentation..... | 73 |
| Figure 4.2.2: Yeast cell growth and biomass accumulation. | 74 |
| Figure 4.2.3: Yeast sugar consumption..... | 76 |
| Figure 4.2.4: The pathways involved in carbohydrate metabolism differentially expressed in <i>S. cerevisiae</i> between fermentation conditions.. | 78 |
| Figure 4.3.1: Yeast nitrogen utilization..... | 91 |
| Figure 4.3.2: Distribution of amino acids in unfermented Icewine juice..... | 91 |
| Figure 4.3.3: Distribution of amino acids in Icewine juice 168 h post-inoculation..... | 93 |
| Figure 4.3.4: Distribution of amino acids in diluted Icewine juice 168 h post-inoculation..... | 94 |
| Figure 4.3.5: The central pathways of nitrogen metabolism in <i>S. cerevisiae</i> .. | 101 |
| Figure 4.3.6: The pathways of branched chain amino acid metabolism in <i>S. cerevisiae</i> | 103 |
| Figure 4.4.1: Yeast glycerol production..... | 116 |
| Figure 4.4.2: Probe specificity for <i>STL1</i> validation Northern blots..... | 121 |
| Figure 4.4.3: Expression of <i>STL1</i> during fermentation..... | 121 |
| Figure 4.5.1: Yeast acetaldehyde and acetic acid production. | 127 |
| Figure 4.5.2: The central pathway of fatty acid β -oxidation in <i>S. cerevisiae</i> .. | 134 |
| Figure 4.5.3: <i>ACS1</i> and <i>ACS2</i> probe specificity for validation Northern blots..... | 137 |
| Figure 4.5.4: Expression of <i>ACS1</i> and <i>ACS2</i> during fermentation. | 139 |
| Figure 4.5.5: Expression of <i>ALD3</i> during fermentation..... | 140 |

1. INTRODUCTION

1.1. INTRODUCTION TO THE PROBLEM

Icewine is an intensely sweet specialty dessert wine fermented from the juice of grapes frozen naturally on the vine. Canada is now globally the leading producer by volume of Icewine as production of Canadian Icewines has increased considerably over recent years from 364,635 L in 2000 to 1,177,484 L in 2006 (Vintners Quality Alliance Ontario, 2001 and 2007). However, Icewine production offers many challenges to winemakers such as sluggish fermentation rates, difficulties in reaching 10-11% (v/v) ethanol in the wine before the yeast stop fermenting and controlling volatile acidity in the form of acetic acid (Kontkanen *et al.*, 2004). These fermentation difficulties result from the high concentrations of naturally concentrated soluble solids in the juice, like sugar, that place commercial wine yeast under extreme hyperosmotic stress as they ferment the juice into wine. Although the response of laboratory yeast strains to hyperosmotic stress in an aerobic environment is well understood (reviewed in Hohmann, 2002), there is still limited information on how wine yeast strains respond to stress especially within an oenological environment (reviewed in Pretorius, 2000). Wine yeast strains display physiological properties that are distinct from laboratory strains and these differences can impact the organoleptic properties of wine (reviewed in Remize *et al.*, 1999; Hauser *et al.*, 2001; Rossignol *et al.*, 2003). Although wine yeast have been selected for their oenological properties, there is still much we do not understand about how they differ from laboratory strains, such as their responses to hyperosmotic stress during Icewine fermentation. Also, since Icewine fermentations typically take weeks to months to complete, the initial response by wine yeast to hyperosmotic stress may differ

considerably from their adapted response throughout the fermentation of Icewine juice in addition to differing from the well-characterized initial responses of laboratory strains (reviewed in Hohmann, 2002). Wine yeast responses during stressful fermentations are not well characterized at the molecular level. This lack of understanding is a problem because this stress response leads to production of metabolites that ultimately affects wine quality (Kontkanen *et al.*, 2004; Pigeau and Inglis, 2005; Pigeau *et al.*, 2007) and our limited understanding of the response limits our ability to select and/or design strains better suited to these fermentations.

1.2. OBJECTIVES

The fundamental goal of this project was to gain a more complete understanding of how wine yeast metabolism adapts in response to prolonged exposure to hyperosmotic stress, especially to high concentrations of fermentable sugars found in Icewine juice and how these responses affect metabolite production such that this information could be used in future projects to select yeast best suited for Icewine production. Therefore, the focus of my project was two-fold. Firstly, microarray technology was to be used to identify yeast genes induced or repressed during Icewine fermentation relative to table wine fermentation by comparing the genome-wide expression between these two conditions during the first five days of fermentation. Secondly, differences in gene expression patterns within entire metabolic pathways were to be correlated to yeast cell growth, sugar and nitrogen usage and metabolites produced by the yeast under the two conditions to understand how and why wine yeast respond differently during Icewine fermentation.

1.3. EXPERIMENTAL DESIGN

Differences in gene expression at the genomic level of the commercial wine yeast *Saccharomyces cerevisiae* K1-V1116 fermenting 40°Brix Icewine and 20°Brix diluted juice during the first 5 days of fermentation were compared using cDNA microarrays. Following data analysis, significant changes in gene expression between the two fermentation conditions were interpreted as fold gene induction or repression. These genes were further categorized based on their biological function or participation within specific metabolic pathways using a statistical datamining software package integrated with the *Saccharomyces* genome database (Ashburner *et al.*, 2000; Dwight *et al.*, 2002). The expression of a small subset of genes differentially expressed between fermentation conditions was verified using Northern analysis. To relate the differences in gene expression between juice conditions to juice components consumed during fermentation or metabolites produced by the yeast during fermentation, enzyme assays and HPLC were used to measure metabolite levels throughout the fermentations.

2. LITERATURE REVIEW

2.1. INTRODUCTION TO ICEWINE: PRODUCTION AND CHALLENGES

Icewine is an intensely sweet dessert wine fermented from the juice of grapes frozen naturally on the vine. The first Icewines originated in Germany during the mid-1800's when grapes froze on the vine before harvest thereby producing wines with intense aromas and flavours (Schreiner, 2001). By allowing grapes to freeze on the vine past their normal fall harvest, the berries are subjected to multiple freeze-thaw cycles leading to mechanical and biochemical changes in the berry tissue. Over time, these changes likely contribute to the unique chemical profile of Icewine juice in comparison to grape juice produced from grapes harvested past their normal fall harvest. Today, the successful commercial development of Canadian Icewine into a unique, specialty dessert wine has securely established Canada as a world leader in Icewine production.

Production of Canadian Icewines has increased considerably over recent years (Vintners Quality Alliance Ontario, 2001 and 2007) and as a result, Canada has gained international recognition as a premium wine producing country. Since Icewine is considered to have a key role in the ongoing commercial and economic success of the Canadian wine industry, it is essential that strict standards for Icewine production are met to ensure continued wine quality (Vintners Quality Alliance, 1999).

The Canadian wine industry is regulated by the Vintners Quality Alliance (VQA). Both grape growers and winemakers must follow strict rules and regulations implemented by the VQA covering all aspects of Icewine production, from harvest to finished wine (Vintners Quality Alliance, 1999). Of these regulations, temperature of the harvest is critical to the production of an authentic Icewine. The entire harvesting and

pressing process must be carried out at cold temperatures (-8°C or below) with harvests typically occurring once temperatures reach -11°C . These temperatures must be maintained throughout the pressing process without artificial refrigeration to ensure that the frozen water of the berries does not thaw and dilute the juice. As a result, a juice highly concentrated in all soluble solids including sugars, acids and nitrogenous compounds is extracted as the water remains with the grape skins in the form of ice during pressing. According to the VQA regulations, the concentration of soluble solids in Canadian Icewine juice must reach a minimum of 35°Brix following pressing (Vintners Quality Alliance, 1999) with concentrations of solutes in Icewine musts commonly falling within the range of approximately 38 to 42°Brix (Pigeau *et al.*, 2007).

Icewine production presents many challenges to winemakers as the high levels of naturally concentrated soluble solids like fermentable sugars in the juice place commercial wine yeast under extreme hyperosmotic stress as they ferment the juice into wine (Kontkanen *et al.*, 2004). In comparison to table wine fermentation, the increased osmotic stress yeast experience throughout the course of Icewine fermentation elicits changes in yeast metabolism, which can contribute to fermentation difficulties (Pigeau and Inglis, 2005; Pigeau *et al.*, 2007). Icewine fermentations are often slow and require lengthy periods of time to reach desired ethanol concentrations, with increasing ethanol levels placing yeast under additional stress as alcoholic fermentation progresses. Typically, yeast consume about half of the available sugar and nitrogen during Icewine fermentation such that fermentation ends while there is still considerable residual sugar present (Kontkanen *et al.*, 2004; Pigeau and Inglis, 2005), imparting the intense sweetness characteristic of this wine. Typical ethanol levels in commercial Icewines are

reported at 10.2 % (v/v) with an average of 214.7 g L⁻¹ residual sugar (Nurgel *et al.*, 2004). Stress induced changes in yeast metabolism can also lead to the production of metabolites that ultimately affect wine quality. Of these metabolites, there are concerns regarding the high levels of acetic acid in Icewine produced by wine yeast as a metabolic response to the high osmotic conditions present during Icewine juice fermentation. Acetic acid at high concentrations can be considered an undesirable organoleptic fault in Icewine as it imparts a vinegar aroma to wine when present above its sensory detection threshold. It is also often associated with wine spoilage.

A recent study by Cliff and Pickering (2006) indicates that acetic acid is only detected in Icewine once it reaches a concentration above 3.185 g L⁻¹, whereas the allowable VQA limit of 2.1 g L⁻¹ acetic acid is well below this detection threshold. Furthermore, the average level of acetic acid reported in a survey of 50 Canadian Icewines was only 1.3 g L⁻¹ with a range of 0.5 g L⁻¹ to 2.3 g L⁻¹, with some wines exceeding the allowable limit (Nurgel *et al.*, 2004). So, there is a concern with understanding and controlling volatile acidity during Icewine fermentations in terms of overall wine quality. Since acetic acid can react with ethanol to form ethyl acetate, which is also considered an oxidation fault described as smelling like nail polish remover, high levels of acetic acid may lead to high levels of ethyl acetate. In fact, in the same survey of 50 commercial Icewines, ethyl acetate levels were reported on average to be 0.24 g L⁻¹ (Nurgel *et al.*, 2004), which is very close to (and exceeds) the reported sensory detection threshold for ethyl acetate in Icewine at 0.198 g L⁻¹ (Cliff and Pickering, 2006). In addition to concerns of high acetic acid concentrations leading to high ethyl acetate levels, it has also been reported that wines made from Icewine juice that is very

concentrated (above 42°Brix) may lead to levels of acetic acid that surpass allowable limits (Pigeau *et al.*, 2007). Since Icewine quality is key to its success in the global marketplace, the fundamental goal of this research is to further understand how wine yeast respond and adapt to prolonged hyperosmotic stress during Icewine fermentation and how these responses may affect Icewine quality.

2.2. RESPONDING TO HYPEROSMOTIC STRESS AND OSMOADAPTATION IN *S. CEREVISIAE*

2.2.1. Hyperosmotic stress: cellular effects, consequences and responses

Yeast live in a continually changing environment where the extracellular water activity can fluctuate considerably over the lifetime of the cell. When yeast cells are exposed to hyperosmotic conditions, the immediate effect is the loss of cytosolic water by osmosis driven by the difference in osmotic pressure across the cell membrane. Loss of cytosolic water ultimately leads to changes in cell volume and shape, loss of turgor pressure, compression of the cytoplasm, oxidative damage and changes in metabolism (Blomberg and Adler, 1992; Varela and Mager, 1996; Billi and Potts, 2002). Thus, maintaining intracellular water homeostasis is fundamental for survival.

Osmoadaptation is a cellular survival response used by yeast to adjust to high external osmolarity by re-establishing the osmotic gradient across the plasma membrane. This response is associated with the increased intracellular accumulation of compatible osmolytes to counteract the efflux of water, thereby restoring cell volume and turgor pressure. Glycerol is the main internal osmolyte produced by *S. cerevisiae* under hyperosmotic conditions (reviewed in Nevoigt and Stahl, 1997).

Adaptation to increased osmolarity is a dynamic process dependent on sensing osmotic changes and relaying these changes to initiate appropriate cellular responses

aimed at maintaining normal biological activities in the cell. The principal cellular response in yeast involved in adaptation to hyperosmotic stress is the high osmolarity glycerol (HOG) response (Brewster *et al.*, 1993). Yeast are also capable of invoking a general stress response which also has an important role in cell survival under a range of stress conditions, including exposure to hyperosmolarity (reviewed in Hohmann, 2002). However, the length of the osmoadaptive phase can vary and is dependent on the type and concentration of stress-inducing solute, yeast strain and duration of exposure to osmotic stress (Blomberg *et al.*, 1988). During Icewine juice fermentation, the stresses placed on wine yeast are persistent and continually changing.

2.2.2. Signaling pathways involved in the hyperosmotic stress response

2.2.2.1. The HOG response

Adaptation to hyperosmotic stress in *S. cerevisiae* is controlled, in part, by an osmosensing and signaling system called the HOG (High Osmolarity Glycerol) MAPK (Mitogen-activated protein kinase) pathway. Evidence of a central MAPK-dependent phosphorelay transduction pathway in *S. cerevisiae*, regulating osmotically induced physiological changes in yeast subjected to hyperosmotic stress, was provided by Brewster *et al.* (1993) using osmosensitive mutants. MAPK signal transduction pathways are highly conserved among eukaryotes and participate in a number of central cellular processes including growth, morphogenesis and stress response. In addition to the HOG pathway, there are five other distinct MAPK pathways in *S. cerevisiae*, involved in sporulation, cell integrity, vegetative growth, pseudohyphal development and mating (reviewed in Hohmann, 2002).

The signaling mechanism of the HOG pathway is based on the sequential phosphorylation of three levels of protein kinases: a MAP kinase kinase kinase (MAPKKK), a MAP kinase kinase (MAPKK) and a MAP kinase (MAPK). In general, the MAPKKK activates the MAPKK via phosphorylation of N-terminal serine and threonine residues residing in the kinase domain. The MAPKK subsequently phosphorylates a threonine or serine residue and a tyrosine residue located within the activation loop of the MAPK catalytic domain. Dual phosphorylation of the threonine/serine and tyrosine residues is required for MAPK activation and subsequent translocation into the nucleus where it then activates downstream transcriptional regulatory proteins (Figure 2.1) (reviewed in Hohmann, 2002). These kinases appear to be activated via osmosensors where changes in extracellular osmolarity generate an internal signal that is transmitted to the HOG MAP kinase cascade. In *S. cerevisiae*, two distinct transmembrane protein osmosensors, Sln1p (Ostrander and Gorman, 1999) and Sho1p (Reiser *et al.*, 2000), have been implicated in sensing osmotic stress. Mutations in the genes encoding for Sln1p (Maeda *et al.*, 1994) and Sho1p (Posas and Saito 1997) affect the activity of the HOG pathway thereby placing the osmosensors upstream of all other HOG pathway components.

Although the mechanism of sensing extracellular osmolarity has not been fully elucidated, observations by Tamás *et al.* (2000) suggest that the signal perceived by the osmosensors results from a change in membrane tension due to turgor stress imposed on the cell rather than a change in extracellular water activity. In yeast, both branches of the HOG pathway were reported to be differentially activated in response to different degrees of salt-induced hyperosmotic stress (O'Rourke and Herskowitz, 2004). Following

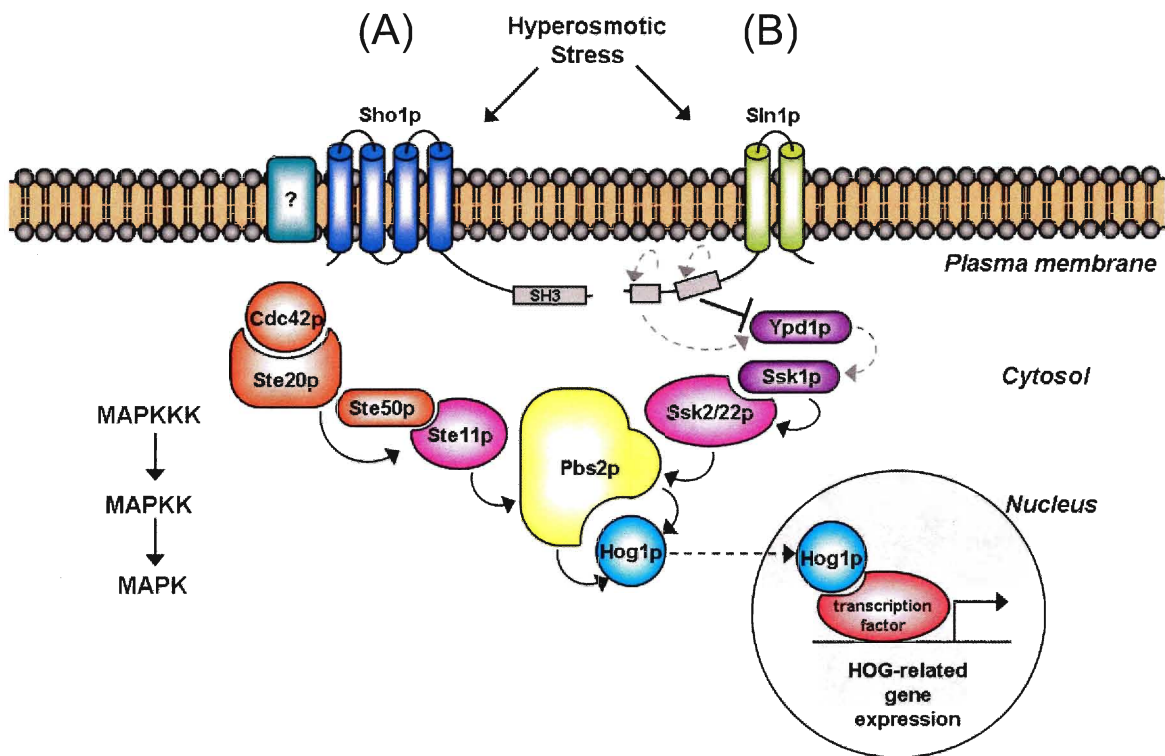


Figure 2.1: The HOG pathway in *S. cerevisiae*. The major components of the high osmolarity glycerol (HOG) pathway including the Sho1p (A) and Sln1p (B) branches are outlined. Solid black arrows indicate associative and phosphorylation events between protein complexes and MAP kinases required for signal transmission under hyperosmotic conditions. Dashed grey arrows indicate phosphotransfer events under normal growth conditions. Adapted from Hohmann (2002) and O'Rourke *et al.* (2002).

transcriptional analysis using microarray technology, O'Rourke and Herskowitz (2004) found that genes associated with the Sln1-branch, but not Sho1p, were induced in yeast exposed to low osmolarity (up to 0.125 M KCl) whereas transcription of genes associated with both branches of the pathways were induced under moderately high hyperosmotic stress (>0.5 M KCl). At higher osmolarities, genes involved in both branches of the HOG pathway as well as the general stress response pathway were involved in the response to hyperosmotic stress.

Central to the HOG pathway are three MAPKKK (Ssk2p, Ssk22p and Ste11p), a MAPKK (Pbs2p) and a MAPK (Hog1p) (reviewed in Hohmann, 2002). The HOG pathway is divided into two independently operating branches, the Sho1p and Sln1p branches, upstream of both Pbs2p and Hog1p (Figure 2.1). Blocking either branch of the pathway still permits phosphorylation and activation of Hog1p upon osmotic shock (Maeda *et al.*, 1995). However, osmosensors Sho1p and Sln1p have very different functional domains and therefore activate different downstream effector proteins within the HOG pathway.

Sho1p contains four transmembrane domains and a cytosolic C-terminal SH3 linker domain (Figure 2.1) (Posas and Saito, 1997). Results from a study by Raitt *et al.* (2000) suggest that Sho1p might not sense osmolarity changes directly, but may instead provide a membrane-bound docking site for Pbs2p via the SH3 linker domain, indicating the possible involvement of a yet unidentified sensor protein or an alternate mechanism for signal transmission through this branch of the HOG pathway. HOG pathway activation by the Sho1p branch relies on the formation of a transient multiprotein osmosignaling complex at the plasma membrane following osmotic shock consisting of

Sho1p, Cdc42p (a GTPase), Ste20p (p21-activated kinase), Ste50p (a SAM domain containing protein), Ste11p (MAPKKK) and Pbs2p (MAPKK) (Reiser *et al.*, 2000; reviewed in O'Rourke *et al.* 2002). The Sho1p branch of the osmosensing HOG pathway and the pseudohyphal development pathway share several MAP kinase components including Sho1p itself (reviewed in Hohmann, 2002). Cdc42p plays a potential role in recruiting the Pbs2p complex to the plasma membrane, as the localization and the physical interaction of Pbs2p with the Sho1p SH3 domain is required for activation of this HOG pathway branch (Reiser *et al.*, 2000). Within the complex, Ste11p is activated by Ste20p (Gustin *et al.*, 1998). Osmotress induced activation of Ste11p is dependent on its association with a putative protein cofactor Ste50p (Posas *et al.*, 1998; Jansen *et al.*, 2001). Activated Ste11p phosphorylates Pbs2p and in turn, Pbs2p phosphorylates cytosolic Hog1p (MAPK) (Posas and Saito, 1998). Phosphorylated Hog1p then translocates into the nucleus (Reiser *et al.*, 1999; Ferrigno *et al.*, 1998). Once in the nucleus, Hog1p regulates the expression of several HOG pathway targeted osmoresponsive genes mediated by its association with several transcriptional regulators.

The Sln1p branch functions as a negative regulator of the HOG pathway under normal growth conditions. Sln1p is comprised of two N-terminal transmembrane domains separated by a loop, an intracellular linker region, a histidine kinase domain and a receiver domain (Figure 2.1) (Ota and Varshavsky, 1993). The transmembrane domains of Sln1p likely function as osmosensors (Ostrander and Gorman, 1999), while the histidine kinase and receiver domains together form a two-component system responsible for relaying signals to two other downstream effector proteins Ypd1p and Ssk1p (West and Stock, 2001). Sln1p-Ypd1p-Ssk1p forms a compact phosphorelay system that

regulates the HOG pathway (Posas *et al.*, 1996). Under normal growth conditions, Sln1p is constitutively activated by autophosphorylation of an internal histidine. The phosphate is intramolecularly transferred to an aspartic acid residue within the receiver domain and in turn, the phosphate is then passed to a histidine residue in Ypd1p. The response regulator, Ssk1p, is the final phosphoacceptor protein in this system. Subsequent transfer of the phosphate group from Ypd1p to Ssk1p deactivates Ssk1p (Posas *et al.*, 1996) consequently preventing activation of the downstream MAPKKs Ssk2p and Ssk22p within the Sln1 branch of the HOG pathway (Posas *et al.*, 1998). Deletion of *SLN1* and/or *YPD1* is lethal due to HOG pathway overactivation (Maeda *et al.*, 1994). Exposure to hyperosmotic stress inhibits the histidine kinase activity of Sln1p thereby disrupting the phosphorelay cascade and thus enabling the binding of dephosphorylated Ssk1p to Ssk2p/Ssk22p (Maeda *et al.*, 1995). Association with dephosphorylated Ssk1p triggers autophosphorylation of Ssk2p/Ssk22p (Posas and Saito, 1998), followed by phosphorylation of Pbs2p and ultimately the activation of Hog1p.

Under hyperosmotic stress, the Sho1p and Sln1p branches of the HOG pathway merge and the activated MAPKK Pbs2p that is shared between branches phosphorylates the cytosolic MAPK Hog1p (Posas and Saito, 1998). Phosphorylated Hog1p then translocates into the nucleus (Figure 2.1) (Ferrigno *et al.*, 1998; Reiser *et al.*, 1999). Once in the nucleus, Hog1p regulates the expression of several osmoreponsive genes mediated by its association with several transcription factors like Hot1p. In turn, the expression of a small subset of HOG pathway targeted genes, including the upregulation of genes encoding for key enzymes involved in glycerol synthesis (*GPD1* and *GPP2*) is affected (Rep *et al.*, 1999; Rep *et al.*, 2000). Yeast cells compensate for the loss of cytosolic water

by accumulating an internal osmolyte, glycerol, through increased biosynthesis when placed under high extracellular osmolarity conditions (Blomberg and Adler, 1989). Mutations in or deletions of Pbs2p or Hog1p cause osmosensitivity in yeast, in addition to reducing intracellular glycerol accumulation (Brewster *et al.*, 1993).

Upon adaptation to hyperosmotic stress, three serine/threonine phosphatases (Ptc1p, Ptc2p and Ptc3p) and two tyrosine phosphatases (Ptp2p and Ptp3p) are upregulated. These phosphatases function to dephosphorylate Hog1p and/or the other MAP kinases in order to deactivate the HOG pathway (Warmka *et al.*, 2001; Jacoby *et al.*, 1997; Wurgler-Murphy *et al.*, 1997). HOG pathway deactivation is necessary once yeast cells return to normal growth conditions since constitutive Hog1p activation in yeast is lethal (Maeda *et al.*, 1993; Jacoby *et al.*, 1997).

2.2.2.2. *The general stress response*

Yeast can invoke an additional cellular response upon exposure to a broad spectrum of stress conditions including heat shock, nutrient starvation together with oxidative and osmotic stress (reviewed in Hohmann, 2002). Similar to the HOG response, the general stress response ultimately affects the expression of a number of genes which have key roles in protecting yeast cells exposed to multiple stress conditions. Of these general stress-responsive genes, a considerable subset contain stress response elements (STREs) within their promoter regions and their expression is controlled by the transcription factors Msn2p and Msn4p (Martinez-Pastor *et al.*, 1996). Both Msn2p and Msn4p contain DNA-binding zinc finger domains predicted to interact with the core CCCCT sequence within the STRE promoter regions (Martinez-Pastor *et al.*, 1996). Control of STRE targeted gene expression is regulated by cellular localization of

Msn2p/Msn4p through protein kinase A (PKA) activity (Görner *et al.*, 1998). Exposure to various stress conditions like high extracellular osmolarity results in the nuclear translocation of Msn2p and Msn4p, causing the upregulation of stress responsive genes, whereas PKA activity mediates the translocation of Msn2p/Msn4p from the nucleus back to the cytosol (Görner *et al.*, 1998). There is evidence supporting an overlap between the HOG pathway and the general stress response when yeast are subjected to hyperosmotic stress (Schüller *et al.*, 1994; Rep *et al.*, 2000; O'Rourke and Herskowitz, 2004). The simultaneous transcriptional activation of some osmoresponsive genes by both HOG-specific stress signaling and STRE-dependent promoter control may be necessary to ensure an effective stress response.

2.2.2.3. Immediate response to hyperosmotic stress

It has been previously shown that Icewine juice contains concentrated solutes in addition to sugar that increase stress on wine yeast fermenting this juice (Pigeau and Inglis, 2005). Although the concentration of salt present in Icewine juice is largely unknown and has not been measured to date, the presence of salts in this juice could potentially intensify the overall hyperosmotic stress wine yeast face during Icewine fermentation, inducing not only the HOG pathway and the general stress response but also an immediate response.

The uptake and export of ions play an important role during osmoadaptation in essentially all types of organisms (reviewed in Hohmann, 2002). Yeast cells specifically control intracellular sodium and potassium ion homeostasis by actively exporting Na^+ and accumulating K^+ (reviewed in Serrano and Rodriguez-Navarro, 2001; reviewed in Hohmann, 2002). Na^+ and K^+ transport is mediated either by ATP-dependent active

transport or by H^+ antiport systems, which are dependant upon the proton gradients established and maintained by ATPases across both the cytoplasmic and vacuolar membranes (reviewed in Serrano and Rodriguez-Navarro, 2001; reviewed in Hohmann, 2002). Although these mechanisms are particularly important for maintaining ion homeostasis in yeast exposed to salt stress, several ion transporters also have roles in osmotic adaptation and/or are regulated by osmotic stress (reviewed in Hohmann, 2002). Nha1p is a yeast plasma membrane K^+/H^+ and Na^+/H^+ antiporter that increases Na^+ efflux in yeast cells exposed to sodium chloride stress (Proft and Struhl, 2004) and decreases K^+ efflux from cells subjected to sorbitol stress (Kinclova-Zimmermannova and Sychrova, 2006). Maintenance of intracellular ion balance via the expression of *NHA1* may have an integral role in facilitating a yeast cell's ability to mount an immediate response to hyperosmotic stress as activation of *NHA1* may be HOG-dependent (Proft and Struhl, 2004; Kinclova-Zimmermannova and Sychrova, 2006).

2.2.3. Glycerol metabolism in *S. cerevisiae*

Glycerol is a major by-product of yeast alcoholic fermentation, in addition to serving as an internal osmolyte during hyperosmotic stress. During typical alcoholic fermentation, approximately 5% of sugars present in grape juice are not completely converted to ethanol and carbon dioxide, but are diverted towards glycerol biosynthesis (reviewed in Scanes *et al.*, 1998). During table wine fermentations, glycerol biosynthesis in yeast primarily serves to maintain balance between redox equivalents by reoxidizing NADH produced during biomass formation (van Dijken and Scheffers, 1986). Glycerol contributes to the sensory qualities of table wine including taste, viscosity and mouthfeel (reviewed in Scanes *et al.*, 1998). The threshold for perceived sweetness and mouthfeel in

table wine due to the presence of glycerol is 5.2 g L⁻¹ and 25.8 g L⁻¹, respectively (Noble and Bursick, 1984). A survey of Canadian Icewines reported an average glycerol concentration of 12.4 g L⁻¹ (Nurgel *et al.*, 2004). The contribution of glycerol to the overall perceived sweetness and mouthfeel in Icewine at this concentration is not clear but it is part of an ongoing investigation (Nurgel and Pickering, 2005).

Three main purposes for glycerol production during wine fermentation have been established, two of which are involved in maintaining cytosolic redox balance during the early stages of fermentation and the third is in response to hyperosmotic stress. Glycerol can be synthesized by commercially available, aerobically grown lyophilized wine yeast strains when introduced to an anaerobic grape juice environment. Glycerol is chiefly produced to reestablish the balance in cytosolic redox equivalents within the nicotinamide cofactor system by oxidizing the additional NADH generated during yeast biomass accumulation under anaerobic conditions like those found during table wine fermentation (van Dijken and Scheffers, 1986). Glycerol synthesis also permits yeast to fulfill the demand for oxidized cofactor, NAD⁺, in response to the build up of NADH generated in glycolysis when the enzymes associated with alcoholic fermentation that would normally oxidize NADH to NAD⁺ are not present in sufficient concentrations to cope with the initial stages of sugar assimilation upon inoculation (reviewed in Bisson, 1993).

2.2.3.1. Glycerol synthesis during hyperosmotic stress

Upon exposure to hyperosmotic stress, water is recruited from the vacuole into the cytoplasm to partially compensate for the sudden loss of cytosolic water by osmosis and the subsequent increase in intracellular osmolarity (Latterich and Watson, 1993;

Mager and Siderius, 2002). This mechanism permits cell survival until yeast begin to accumulate glycerol to adjust to changes in external water activities. Studies involving laboratory yeast strains grown under salt stress have shown that glycerol functions as an internal osmolyte to counterbalance the osmotic pressure placed on cells, and to counteract water loss and help cells resume growth (Blomberg and Adler, 1989; Blomberg and Adler, 1992; Brewster *et al.*, 1993; Blomberg, 2000). Net glycerol accumulation in yeast under hyperosmotic stress may be dependent on a series of events: increased production, decreased catabolism, increased retention by the plasma membrane and/or increased uptake of extracellular glycerol (reviewed in Nevoigt and Stahl, 1997).

In *S. cerevisiae*, glycerol is synthesized from the glycolytic intermediate dihydroxyacetone phosphate (DHAP) in two enzymatic reactions. The first reaction requires the reduction of DHAP to glycerol-3-phosphate along with the simultaneous oxidization of NADH to NAD⁺ (Figure 2.2). This rate-limiting step (Remize *et al.*, 2001) in glycerol synthesis is catalyzed by a NAD⁺-dependent glycerol-3-phosphate dehydrogenase encoded by the highly homologous genes *GPD1* and *GPD2* (Larsson *et al.*, 1993; Eriksson *et al.*, 1995). The second reaction is completed following the dephosphorylation of glycerol-3-phosphate to glycerol by glycerol-3-phosphatases (*GPP1* and *GPP2*) (Påhlman *et al.*, 2001). This pathway appears to be the sole route identified to date for glycerol production in *S. cerevisiae*.

The genes encoding for the isoforms of glycerol-3-phosphate dehydrogenase and glycerol-3-phosphatase are differentially expressed and appear to be controlled by different physiological triggers. *GPD1* expression is upregulated upon the exposure of cells to hyperosmotic stress and is considered to be the major isoform responsible for

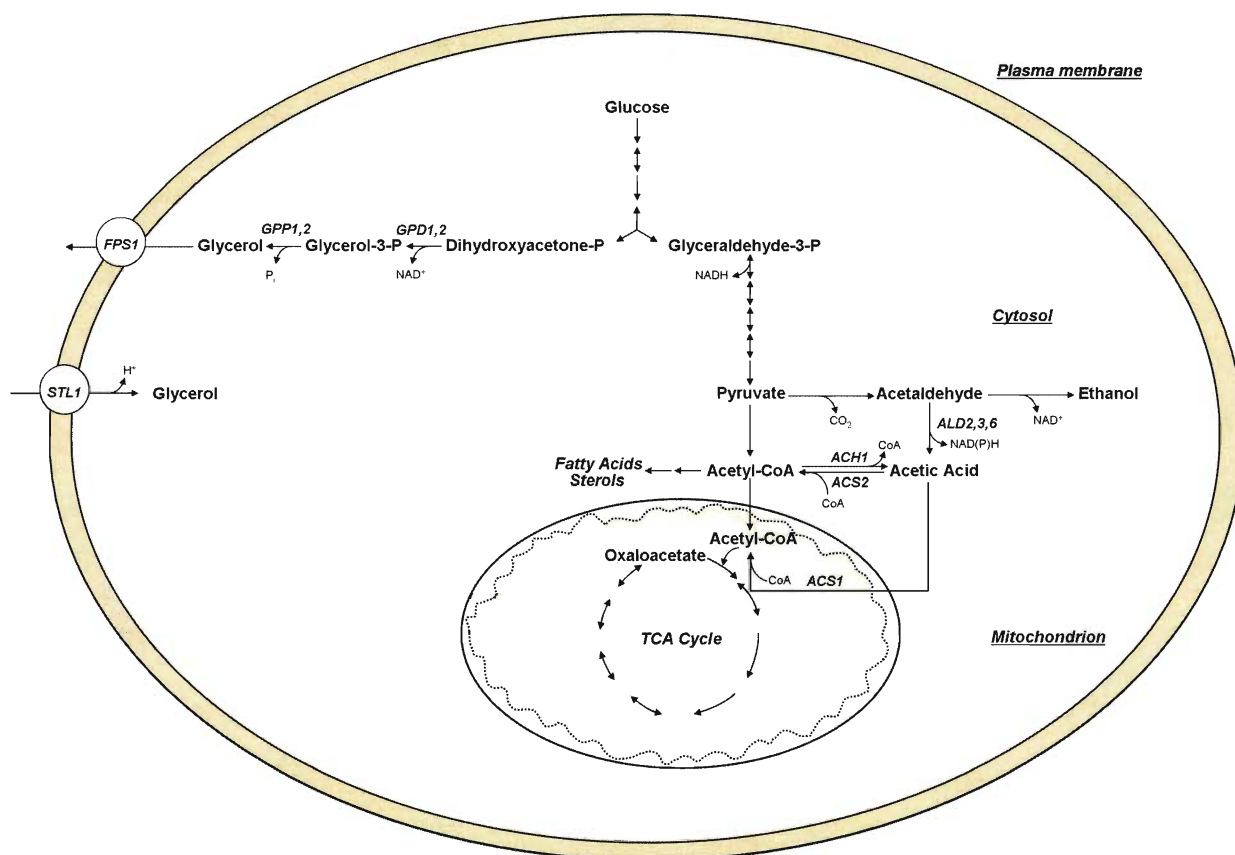


Figure 2.2: Glycerol and acetic acid metabolic pathways in *S. cerevisiae*. The pathways of glycerol and acetic acid metabolism and the genes encoding for enzymes involved in these pathways are illustrated. Metabolic intermediates and cofactors are also depicted. *GPD1,2*- Glycerol-3-phosphate dehydrogenase; *GPP1,2*- Glycerol-3-phosphatase; *FPS1*- glyceroporin channel; *STL1*- glycerol/H⁺ symporter; *ALD2,3,6*- aldehyde dehydrogenase; *ACS1,2*- acetyl-CoA synthetase and *ACH1*- acetyl-CoA hydrolase.

glycerol production during the hyperosmotic response by yeast (Larsson *et al.*, 1993; Albertyn *et al.*, 1994a). In comparison, *GPD2* expression is slightly repressed under hyperosmotic conditions but is upregulated in the absence of oxygen (Ansell *et al.*, 1997). Expression of *GPD2* functions to maintain intracellular redox balance under anaerobic conditions (Eriksson *et al.*, 1995; Ansell *et al.*, 1997). The *GPD1* promoter binds Hog1p (Alepuz *et al.*, 2001), yet the HOG pathway was found to exert limited control of *GPD1* expression during table wine fermentations (Remize *et al.*, 2003). The *GPD1* promoter also contains three STREs (Eriksson *et al.*, 2000) indicating that expression is regulated by both the HOG pathway and the general stress response. In contrast, *GPD2* expression is thought to be regulated by the redox state of the cell (Ansell *et al.*, 1997).

GPP1 and *GPP2* are both induced by hyperosmotic conditions such that salt stress induced expression was found to be dependent on HOG pathway activation (Norbeck *et al.*, 1996; Rep *et al.*, 2000; Pählman *et al.*, 2001). *GPP2* expression is more responsive in yeast exposed to saline stress (Hirayama *et al.*, 1995), whereas *GPP1* expression is found to be less responsive (Norbeck *et al.*, 1996). However, *GPP1* expression exhibits transient upregulation when yeast cells are transferred to anaerobic conditions, while *GPP2* expression remains unaffected (Pählman *et al.*, 2001). Under hyperosmotic conditions, expression of *GPP2* is independent of the STRE binding general stress response transcription factors Msn2p and Msn4p (Rep *et al.*, 1999) and PKA activity (Norbeck and Blomberg, 2000), while *GPP1* expression is diminished in yeast strains with low PKA activity (Pählman *et al.*, 2001).

The combination of Gpd1p-Gpp2p activity is proposed to be mainly involved in glycerol production during osmotic stress, whereas the Gpd2p-Gpp1p combination is

associated with balancing the NAD^+/NADH cofactor system under anaerobic conditions (Albertyn *et al.*, 1994b; Ansell *et al.*, 1997; Pålman *et al.*, 2001). Wine yeast face both anaerobic and osmotic stress during fermentation and either of these factors may control the expression of *GPD* and *GPP* genes. However, the elevated production of glycerol by wine yeast during Icewine fermentation corresponds to an increase of *GPD1* expression (Pigeau and Inglis, 2005) but not of *GPD2* (Pigeau and Inglis, 2007).

Glycerol can be used as a sole carbon source under aerobic conditions. A cytosolic glycerol kinase encoded by *GUT1* (Pavlik *et al.*, 1993) phosphorylates glycerol to form glycerol-3-phosphate and in turn is oxidized to DHAP, which then enters the glycolytic pathway. This reaction is catalyzed by a mitochondrial FAD^+ -dependent glycerol-3-phosphate dehydrogenase encoded by *GUT2* (Rønnow and Kielland-Brandt, 1993). Expression of both *GUT1* and *GUT2* is subject to glucose repression and does not appear to be induced by osmotic stress (reviewed in Hohmann, 2002). A decrease in glycerol catabolism in yeast under hyperosmotic stress may be necessary to maintain intracellular glycerol levels.

2.2.3.2. Glycerol retention and uptake

To ensure that the glycerol produced in *S. cerevisiae* upon exposure to hyperosmotic stress is accumulated to offset the osmotic pressure exerted on the plasma membrane, yeast must simultaneously rely on increased glycerol biosynthesis and decreased glycerol export through the plasma membrane. Of the two major glycerol transport systems currently identified in *S. cerevisiae*, one system involves a channel protein belonging to the Major Intrinsic Protein (MIP) family and is encoded by the gene *FPS1* (Van Aelst *et al.*, 1991), as illustrated in Figure 2.2.

The rate of simple diffusion of glycerol through the plasma membrane is low, however membrane permeability to glycerol is reduced even further in osmotically stressed yeast cells (Tamás *et al.*, 1999; Sutherland *et al.*, 1997). There may be a connection between the modification of the fatty acid composition of phospholipids and ergosterol levels in the lipid bilayer of the plasma membrane to reduced permeability to glycerol to regulate transmembrane flux of glycerol (reviewed in Nevoigt and Stahl, 1997; Hohmann, 2002). The main physiological role of Fps1p appears to be exporting glycerol from yeast cells upon hypoosmotic shock or a return to normal growth conditions following hyperosmotic shock, by facilitating passive transport of glycerol across the membrane down a concentration gradient (Van Aelst *et al.*, 1991; Luyten *et al.*, 1995; Sutherland *et al.*, 1997; Oliveira *et al.*, 2003). It has been suggested that Fps1p can transport glycerol in both directions across the membrane (Sutherland *et al.*, 1997; Tamás *et al.*, 1999). However, this channel rapidly closes upon exposure to hyperosmotic stress, thereby controlling the amount of glycerol retained by the cell (Támas *et al.*, 1999). Therefore, it is unlikely that Fps1p would be involved in glycerol uptake in osmotically stressed yeast cells.

It has been shown that deletion mutants lacking Fps1p retain more glycerol than their wild-type counterparts following exposure to salt stress (Luyten *et al.*, 1995). The N-terminal regulatory domain of Fps1p is responsible for controlling channel opening and closing (Támas *et al.*, 1999). Deletion of this domain results in constitutive glycerol export, thereby causing yeast cells to overproduce glycerol in order to counterbalance the loss of cytosolic glycerol (Támas *et al.*, 1999; Remize *et al.*, 2001). Regulation of Fps1p closing upon hyperosmotic stress is still unclear but may require HOG pathway activation

and/or may depend on sensing changes in cytoskeleton architecture (reviewed in Hohmann, 2002).

Osmotolerant yeasts like *Zygosaccharomyces rouxii* actively uptake glycerol against a concentration gradient and it has been suggested that a high affinity glycerol/H⁺ symporter uptake system also exists in *S. cerevisiae* (van Zyl *et al.*, 1990; Lages and Lucas, 1997). Active glycerol transport was first proposed to be mediated in *S. cerevisiae* by the putative multi-membrane spanning glycerol uptake proteins Gup1p and Gup2p (Holst *et al.*, 2000). *GUP1* expression was found to be necessary for growth on nonfermentable carbon sources, specifically glycerol as the sole carbon source, while *GUP2* expression was found to be induced by salt stress (Holst *et al.*, 2000). However, a study by Oliveira and Lucas in 2004 revealed that *GUP1* and *GUP2* transcriptional activity was found to be constitutive and not affected by either glucose repression or salt stress in laboratory strains of *S. cerevisiae*. Also, glycerol uptake was also observed in $\Delta gup1 gup2 gut2$ deletion mutants grown under salt stress, suggesting the existence of another protein involved in active glycerol uptake in *S. cerevisiae* (Neves *et al.*, 2004). *STL1* was first characterized as encoding a putative sugar transporter-like protein (Nelissen *et al.*, 1997) and its expression was found to be strongly but transiently induced by salt stress and required HOG pathway activation (Rep *et al.*, 2000). *STL1* was also one of the most strongly upregulated genes in wine yeast following a two hour exposure to high sugar stress (Erasmus *et al.*, 2003). Recently, Stl1p (Figure 2.2) has been characterized as a glycerol/H⁺ symporter subject to glucose repression in aerobically grown laboratory yeast strains (Ferreira *et al.*, 2005). Active glycerol uptake has not been investigated in osmotically stressed wine yeast strains in the presence of high

concentrations of fermentable sugars. It is not currently known if this is a mechanism that wine yeast utilize during Icewine juice fermentation to combat hyperosmotic stress in order to add to the cytosolic pools of biosynthesized glycerol.

2.2.4. Acetic acid metabolism in *S. cerevisiae*

Fermentation of glucose to ethanol via glycolysis and alcoholic fermentation is a redox neutral process whereby the NADH generated during glycolysis is reoxidized during alcoholic fermentation. Glycerol synthesis is not a redox neutral process so a redox imbalance in the NAD^+/NADH cofactor system is potentially set up while osmotically stressed wine yeast cells are fermenting sugar to ethanol. Yeast are unable to convert reducing equivalents between the NAD^+/NADH and $\text{NADP}^+/\text{NADPH}$ cofactor systems as they lack transhydrogenase activity, so they must rely on alternate metabolic pathways to maintain intracellular redox balance between these cofactor systems (van Dijken and Scheffers, 1986). Acetic acid synthesis has been suggested as a mechanism through which yeast can metabolically reestablish cytosolic redox balance by regenerating NADH from the NAD^+ produced during glycerol biosynthesis in response to salt induced hyperosmotic stress (Blomberg and Adler, 1989; Wang *et al.*, 1998; Navarro-Avino *et al.*, 1999). This pathway involves the oxidation of acetaldehyde to acetic acid catalyzed by an aldehyde dehydrogenase (Figure 2.2).

Of the five aldehyde dehydrogenases known in *S. cerevisiae*, only three are specific to NAD^+ (*ALD2*, 3 and 4) and may solve the redox imbalance during glycerol production. However, the NADP^+ -dependent Ald6p was found to be the main aldehyde dehydrogenase responsible for acetic acid production in laboratory *S. cerevisiae* strains fermenting glucose media (Remize *et al.*, 1999; Eglinton *et al.*, 2002), while Ald5p was

also found to contribute to acetic acid biosynthesis during anaerobic growth on glucose (Saint-Prix *et al.*, 2004). Of these two NADP⁺-dependent isoforms, only *ALD6* was upregulated under salt stress and its expression required HOG pathway activation (Akhtar *et al.*, 1997; Rep *et al.*, 2000). Furthermore, deletion of *ALD6* in wine yeast fermenting synthetic must containing 200 g L⁻¹ of sugar, resulted in a decrease in acetic acid production and an increase in the formation of other metabolites, specifically glycerol, succinate and butanediol (Remize *et al.*, 2000). Although it is clear that Ald6p plays a role in acetic acid production during fermentation, its function, if any, in resolving redox imbalance under these hyperosmotic stress conditions where glycerol is made, is not clear due to its cofactor specificity.

Expression of the highly homologous *ALD2* and *ALD3* isoforms was thought to be both subject to osmotic stress induction and regulated by the general stress transcription factors Msn2p and Msn4p (Martinez-Pastor *et al.*, 1996), but only *ALD3* expression was observed to be responsive to salt stress in laboratory yeast strains (Norbeck and Blomberg, 2000). Ald2p activity has been associated with catalyzing the oxidation of 3-aminopropanal to form β -alanine, which is required for pantothenic acid production (White *et al.*, 2003). The contribution of aldehyde dehydrogenase activity to increased acetic acid levels produced by wine yeast strains in the presence of high concentrations of fermentable sugars during Icewine fermentation is not fully understood. In a study by Pigeau and Inglis (2005, 2007), of all the *ALD* genes, only *ALD3* has been found to be more highly expressed during Icewine fermentation in comparison to that found during a table wine fermentation and this increased expression corresponded to an increase in acetic acid in the wine. Although the cytosolic isoform *ALD6* was expressed

throughout Icewine juice fermentation, its expression was not significantly different when compared to dilute juice fermentation.

To date, it is not clear if Ald3p plays a role in the elevated acetic acid production during Icewine fermentation. Studies by Pigeau and Inglis (2005 and 2007) indicate that of all the aldehyde dehydrogenases, only *ALD3* shows an increased expression in Icewine fermentation above that found during the table wine fermentation. However, two recent studies have shown that deletion of *ALD3* from the yeast genome did not reduce acetic acid production (Cordier *et al.*, 2007; Modig *et al.*, 2007). It was also reported that overexpression of *ALD3* in a laboratory yeast strain did not significantly change the amount of acetic acid produced by cells exposed to salt stress (Modig *et al.*, 2007). It is likely that the expression of *ALD3* plays a role in maintaining redox balance and ultimately contributes to elevated acetic acid levels in Icewine (Pigeau and Inglis, 2005), but enzymatic activity of Ald3p has yet to be reported.

2.2.5. Pyruvate dehydrogenase bypass

Acetic acid is an intermediate metabolite of the pyruvate dehydrogenase (PDH) bypass in *S. cerevisiae* leading to the formation of acetyl-CoA from pyruvate (reviewed in Pronk *et al.*, 1996). In this pathway, acetic acid can be converted to acetyl-CoA by acetyl-CoA synthetase. In yeast, two genes (*ACS1* and *ACS2*) have been identified as encoding for proteins associated with acetyl-CoA synthetase activity (Figure 2.2). *ACS1* is the mitochondrial isoform required for respiratory growth on non-fermentable carbon sources and is subject to glucose repression (Kratzer and Schüller, 1995; Van den Berg *et al.*, 1996). *ACS2* is the cytosolic isoform which is constitutively transcribed and is required for growth on glucose under anaerobic conditions (Van den Berg and Steensma,

1995). Coenzyme A can also be regenerated from the hydrolysis of acetyl-CoA by a cytosolic acetyl-CoA hydrolase (*ACH1*) and is required for growth on acetic acid (Lee *et al.*, 1996). The pool of available cytosolic acetyl-CoA produced by the bypass pathway is utilized for a variety of anabolic reactions, including fatty acid and amino acid biosynthesis (reviewed in Pronk *et al.*, 1996). Akamatsu *et al.* (2000) reported that genes encoding for both acetyl-CoA synthetase isoforms (*ACS1/2*) and the acetyl-CoA hydrolase (*ACH1*) were expressed by an *S. cerevisiae* sake yeast strain fermenting sake mash. In the same study, the expression of a cytosolic NAD⁺-dependent aldehyde dehydrogenase encoded by either *ALD2* or *ALD3* was also observed. However, the authors were unable to design probes specific for these highly homologous isoforms to make a distinction between *ALD2* and *ALD3* expression during sake fermentation. Akamatsu *et al.* (2000) also reported that the overexpression of *ACS2*, but not *ACS1*, resulted in a four-fold reduction in acetic acid production by yeast during small-scale sake fermentations. So, the results from this study suggest that there may be a connection between aldehyde dehydrogenase and acetyl-CoA synthetase activity on acetic acid production by yeast during fermentation. However, expression of *ACS1/2* and *ACH1* during Icewine juice fermentation has not been investigated to date. Acetic acid metabolism in yeast during fermentation is currently not fully understood, but expression of the genes involved in its synthesis along with its degradation may overall contribute to the acetic acid levels produced by wine yeast as they adapt to prolonged exposure to hyperosmotic stress during the fermentation of Icewine juice.

2.3. GENE EXPRESSION PROFILING IN *S. CEREVISIAE* DURING FERMENTATION

Yeast are subjected to continuous fluctuations in their environment and cells must cope with a variety of stress conditions that can eventually affect cell growth and viability. Survival is dependent on rapid adaptation to extracellular stresses and therefore requires substantial genome-wide changes in gene expression to maintain cell integrity and metabolic activity. Under oenological conditions, wine yeast cells are exposed to multiple and concurrent stress conditions during the fermentation of grape juice. Yeast are predominantly challenged with hyperosmotic stress due to high starting sugar concentrations upon inoculation. As the fermentation process proceeds, yeast must contend with ethanol stress and variations in pH, in addition to nutrient depletion and starvation. The overall change in gene expression in conjunction with the corresponding changes in metabolic activity as they respond and adapt to stress during fermentation can lead to the production of compounds that contribute to the sensory properties of the wine. Studying the initial response of wine yeast to a fermentation condition would not take into account these changes in the environment and the impact on gene expression and metabolism.

Commercial *S. cerevisiae* strains currently used for winemaking are closely related to laboratory yeast strains but display unique physiological characteristics allowing them to be more suitable for wine fermentation. The majority of laboratory yeast strains are either haploid or diploid. Wine yeast strains are commonly diploid, aneuploid and/or polyploid and the genome may contain several chromosomal rearrangements (Bidenne *et al.* 1992; reviewed in Pretorius, 2000). An increase in the expression of genes essential for fermentation through an increase in gene copy number or by other genomic reorganizations may play a role in the metabolic properties of wine

yeasts, such as fermentation efficiency, as well as ethanol and osmotic tolerance (reviewed in Pretorius, 2000). There is evidence that the differences in metabolic activity between laboratory and wine yeast strains under fermentative conditions may influence the metabolites produced (Remize *et al.*, 1999).

Genomic expression profiling using microarray technology can uncover the differences in gene expression patterns in yeast under a number of growth and stress conditions. DNA microarrays can measure the relative transcriptional abundance in every gene in the yeast genome. Several microarray experiments have focused on the initial hyperosmotic stress response of laboratory strains exposed to salt stress and will be further discussed in Section 2.3.1. Genome-wide changes in gene expression associated with the adapted stress response to prolonged exposure to sugar stress during Icewine fermentation and its effect on metabolic activity in yeast is yet unclear.

2.3.1. Gene expression of laboratory yeast strains under salt stress

The genome of *S. cerevisiae* was the first eukaryotic genome to be fully sequenced. The earliest array experiments were performed using this organism, and characterized the genes differentially expressed when laboratory yeast strains were grown under differing nutrient conditions associated with carbon source limitation or the diauxic shift (Wodicka *et al.*, 1997; DeRisi *et al.*, 1997; ter Linde *et al.*, 1999). A number of studies have used either salt or sorbitol to determine the changes in gene expression during the hyperosmotic stress response (Posas *et al.*, 2000; Rep *et al.*, 2000; Yale and Bohnert, 2001, Causton *et al.*, 2001). When yeast cells were subjected to a 30-minute exposure to either salt or sorbitol stress, 5% of the yeast genome (286 genes) was found to be induced under hyperosmotic stress, but the expression of only a few genes was

observed to be solute specific. The expression of 10 genes including *ALD3* was upregulated more strongly by salt stress than by sorbitol. In contrast, 26 genes were more strongly upregulated by sorbitol, including *GPD1*, *STL1* and *ALD6* (Rep *et al.*, 2000).

In addition to the number and degree of differential gene expression changes upon exposure to high osmotic stress, the timing of each response can also be affected by the amount of stress. Posas *et al.* (2000) showed the transcriptional response of yeast exposed to salt stress was time and concentration dependent. When placed under a mild (0.4 M) and brief (10 minute) osmotic shock, yeast cells displayed a rapid but transient transcriptional response where approximately 21% of the yeast genome was upregulated at least three-fold. The number of induced genes decreased ten-fold after 20 minutes. However, a higher sodium chloride concentration (0.8 M) resulted in a delayed response, as the number of genes induced after 10 minutes increased two-fold after 20 minutes. In a similar study, Yale and Bohnert (2001) exposed yeast to 1 M sodium chloride for 10, 30 and 90 minutes. It was observed that the changes in transcript abundance greater than two-fold corresponded to an increase in the number of salt-induced genes expressed over time. Genes associated with the initial osmotic stress response after 10 minutes were connected to nucleotide metabolism. The pattern of gene expression after 90 minutes of adaptation to salt stress changed, as the majority of genes induced were associated with energy generation, nitrogen metabolism, fatty acid biosynthesis and the stress response.

Microarray analysis can also illustrate how gene expression patterns can overlap in response to different stresses. Causton *et al.* (2001) performed a time-course expression profile of laboratory yeast strain under multiple conditions including: heat shock, acid and alkali, oxidative, carbon limitation and osmotic stress. Of the 66% of

genes within the yeast genome differentially expressed between the various stresses, most exhibited similar expression patterns over time. The genes significantly upregulated under most environmental stress conditions were related to carbohydrate metabolism, energy generation and cell stress response. Conversely, genes showing a downregulated response were associated with amino acid metabolism, cell wall maintenance and protein synthesis. However, a small subset 370 of genes was uniquely specific for a given environmental stress.

2.3.2. Gene expression profiling in wine yeast

A number of studies have focused exclusively on the genome-wide transcriptional differences between wine yeast strains fermenting synthetic must simulating table wine conditions (Cavalieri *et al.*, 2000; Hauser *et al.*, 2001) or the differences derived when wine yeast are placed under differing nitrogen conditions during fermentation (Backhus *et al.*, 2001; Marks *et al.*, 2003). A comparison of two wine yeast strains revealed significant differences in gene expression as the cells entered early stationary phase during the fermentation of synthetic must (Zuzuarregui *et al.*, 2006). The expression of over 2000 genes was significantly different between strains. Of those genes, 423 showed at least a two-fold difference in mRNA abundance. These differences were primarily related to genes involved in carbohydrate and nitrogen metabolism, in addition to fatty acid and ergosterol synthesis.

In 2003, Rossignol *et al.* profiled the dynamic gene expression changes of a commercially available wine yeast strain throughout the fermentation of synthetic must as cells adapted to changing nutrient and environmental conditions from early exponential to late stationary phase of growth. The greatest number of genes induced or

repressed in cells occurred upon entry into stationary phase, where the expression of over 1000 genes was significantly different when compared to that at the start of the fermentation. The transcriptional response observed corresponded to the changes in nutrient availability, osmotic and ethanol stress.

Erasmus *et al.* (2003) conducted one of the first transcriptional comparisons involving a wine yeast strain of *S. cerevisiae* exposed to high sugar stress. The transcription of 589 genes changed more than two-fold when yeast were exposed for two hours to Riesling grape juice chaptalized with equal amounts of glucose and fructose to a final concentration of 40% (w/v). However, only the initial response to sugar stress was monitored and this response may not relate directly to the metabolic responses that alter wine quality as the yeast continued to ferment for an additional 400 to 500 hours (Kontkanen *et al.*, 2004; Pigeau and Inglis, 2005). One should also note that although Erasmus *et al.* were trying to mimic the composition of Icewine juice in their study, the growth media used did not accurately reflect the high levels of the other naturally concentrated soluble solids in Icewine juice, such as organic acids and salts, which place commercial wine yeast under additional stress (Pigeau and Inglis, 2005). The response of wine yeast as they adapt to prolonged exposure to high osmotic stress throughout Icewine juice fermentation is still unclear and therefore needs to be addressed.

With the use of microarray analysis, our goal in this project was to identify metabolic pathways more or less active during Icewine fermentation, and how these related to the use of nutrients during the fermentation and the production of yeast metabolites involved in redox balance. In addition to more fully understanding acetic

acid and glycerol metabolism under Icewine conditions, overall sugar and nitrogen utilization could be investigated.

2.4. YEAST SUGAR AND NITROGEN METABOLISM

2.4.1. Sugar Utilization

The conversion of grape juice sugars (glucose and fructose) by *S. cerevisiae* to ethanol is of vital importance to wine production (Bisson, 1993). Glucose and fructose are the primary fermentable sugars in grape juice and they are usually present in equal amounts. Under anaerobic conditions, they are cofermented to ethanol and carbon dioxide by *S. cerevisiae* (Fleet and Heard, 1993). The metabolic intermediates of glycolysis and alcoholic fermentation, along with the genes encoding for enzymes associated with these pathways, are depicted in Figure 2.3. Although fructose is used concurrently with glucose during table wine fermentation, *S. cerevisiae* displays a preference for glucose as a carbon source, such that glucose is typically depleted from the juice first, giving rise to a difference between the amount of glucose and fructose consumed during fermentation (Berthels *et al.*, 2004). Previous studies have shown that yeast fermenting Icewine juice only utilize half of the available fermentable sugars present in the juice and are glucophilic (Kontkanen *et al.*, 2004; Pigeau and Inglis, 2005).

In yeast, transporting sugar across the plasma membrane has been proposed as a rate-limiting step of glycolysis by influencing carbon flux through the glycolytic pathway (reviewed in Kruckeberg and Dickinson, 2004). At present, 20 different genes encoding for hexose transport-related proteins have been identified in *S. cerevisiae* (Wieczorke *et al.*, 1999). Expression of a small number of these transporters has been implicated in playing different roles in sugar uptake during wine fermentation (Luyten *et al.*, 2002).

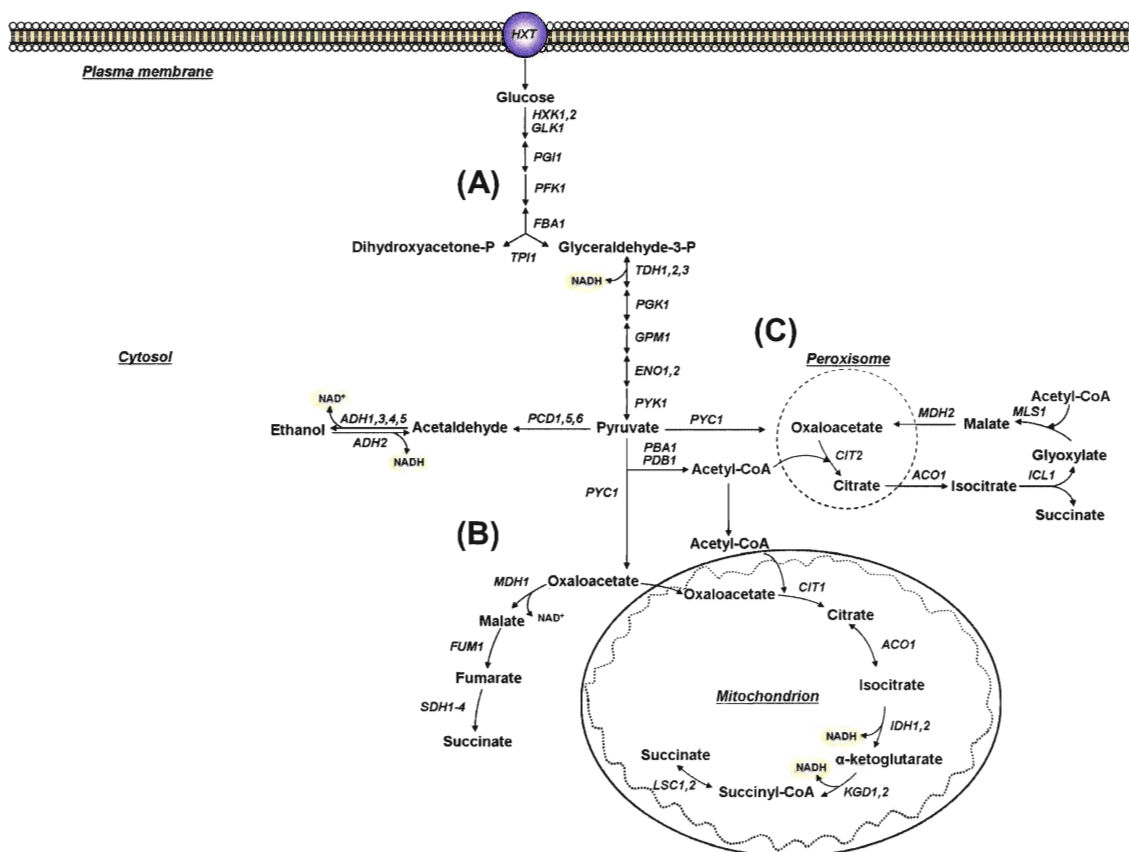


Figure 2.3: Pathways associated with carbohydrate metabolism in *S. cerevisiae*. Illustrated are the metabolic intermediates and the genes associated with (A) glycolysis and alcoholic fermentation, (B) the oxidative and reductive pathways of the tricarboxylic acid (TCA) cycle and (C) the glyoxylate cycle.

There are 18 genes that encode for sugar transporters (*HXT1-17* and *GAL2*) (reviewed in Özcan and Johnston, 1999) and there are two glucose sensors (*SNF3* and *RGT2*) with putative sugar transporter activities (reviewed in Kruckeberg and Dickinson, 2004). Under laboratory conditions, *HXT1-4*, *HXT6* and *HXT7* function as major hexose transporters and deletion of *HXT1-7* abolishes sugar transport altogether in glucose media grown yeast (Reifenberger *et al.*, 1997). Expression of the *HXT* gene family is differentially regulated by the availability of extracellular sugar and all *HXT* gene products have different sugar substrate specificities and affinities. *HXT1* and *HXT3* encode for low affinity transporters expressed in response to glucose levels in excess of 4% (w/v) and were found to play a predominant role in sugar transport in wine yeast fermenting synthetic must (Özcan and Johnston, 1999; Luyten *et al.*, 2002; Perez *et al.*, 2005). *HXT2* and *HXT4* encode for transporters with moderate to low affinity for hexoses. *HXT6* and *HXT7* are associated with high affinity sugar transport in cells under glucose-limited conditions and are repressed by high levels of glucose (Özcan and Johnston, 1999; reviewed in Kruckeberg and Dickinson, 2004). The remaining *HXT* genes (*HXT8-17*) encode for proteins closely related to hexose transporters but it is unclear if they mediate glucose transport, since they are weakly expressed in yeast grown in the presence of differing glucose levels (reviewed in Özcan and Johnston, 1999).

Following transport into the cell, hexoses are phosphorylated by hexokinases upon entry into the glycolytic pathway. *HXX2* is the primary glucose phosphorylating enzyme in yeast growing in conditions of abundant glucose, while the expression of *HXX1* and *GLK1* is required during glucose-limited conditions and is repressed by high glucose conditions (Herrero *et al.*, 1995; Rodriguez *et al.*, 2001). In addition to catalyzing

the first enzymatic reaction of glycolysis, the hexokinases (*HXK1* and *HXK2*) and glucokinase (*GLK1*) may also have key roles in glucose transport in yeast, such that glucose phosphorylation may be required for high affinity glucose uptake (Bisson and Fraenkel, 1983; Bisson and Fraenkel, 1984). *HXK1* and *GLK1* are also responsive to a number of stresses (Gasch *et al.*, 2000). The expression of both genes was induced by salt stress (Posas *et al.*, 2000; Rep *et al.*, 2000; Yale and Bohnert, 2001) and by high sugar stresses (Erasmus *et al.*, 2003). The ability of yeast to ferment hexose sugars is essential for growth and survival under oenological conditions and requires the coupling of glycolysis to alcoholic fermentation to create an energy generating but redox neutral system in the absence of oxygen (van Dijken and Scheffers, 1986).

Pyruvate decarboxylase (*PDC*) and alcohol dehydrogenase (*ADH*) are the key enzymes of alcoholic fermentation. Of the three *PDC* isoforms, *PDC1* is strongly expressed in actively fermenting cells, while conversely, expression of *PDC5* in these same cells cannot be detected (Flikweert *et al.*, 1996; Hohmann and Cederberg, 1990). It has been previously reported that the expression of the minor isoform (*PDC6*) was not involved in glucose fermentation (Hohmann, 1991). However, *PDC6* was strongly upregulated in wine yeast two hours after exposure to high sugar stress but was unaffected by salt stress (Erasmus *et al.*, 2003; Rep *et al.*, 2000). Of the alcohol dehydrogenase isoforms, *ADH1* expression is upregulated in the presence of glucose and is the major cytosolic isoform involved in glucose fermentation (Denis *et al.*, 1983). *ADH2* expression is repressed by glucose and is required for growth on ethanol, while the participation of the other three isoforms (*ADH3-5*) during fermentation has not been fully elucidated (reviewed in Kruckeberg and Dickinson, 2004).

In glycolysis, there is a net production of two ATP molecules by substrate-level phosphorylation and two NADH molecules. Alcoholic fermentation involves the decarboxylation of pyruvate produced during glycolysis to acetaldehyde, which in turn is then reduced to ethanol thereby regenerating NAD^+ . Under aerobic conditions, the pyruvate generated via glycolysis can enter the mitochondria and be converted to oxaloacetate and/or acetyl-CoA, which can enter the tricarboxylic acid cycle (TCA). Alternatively, pyruvate can enter the glyoxylate cycle (reviewed in Krukeberg and Dickinson, 2004). In the presence of fermentable sugars, yeast cells downregulate the expression of genes involved in mitochondrial functions, including respiration and the metabolism of alternate carbon sources such as ethanol and acetate through carbon catabolite repression (Gancedo, 1998).

In *S. cerevisiae*, the tricarboxylic acid (TCA) cycle plays a central role in oxidative growth to provide reducing equivalents used by the mitochondrial respiratory chain to produce ATP (van Dijken and Scheffers, 1986). Under anaerobic conditions, the TCA cycle has been proposed to operate in a noncyclic manner and can be divided into two distinct pathways (Figure 2.3): a cytosolic reductive pathway from oxaloacetate resulting in the formation of NAD^+ or FAD and a mitochondrial oxidative pathway starting from α -ketoglutarate but resulting in the production of NADH (Radler, 1993; Camarasa *et al.*, 2003). It is not known if these pathways operate in yeast during fermentation, especially during an Icewine fermentation, but both pathways can lead to succinate production, which is another important organic acid produced by yeast during grape juice fermentation (Radler, 1993). The glyoxylate cycle typically operates in yeast during respiratory growth, especially when glucose is exhausted and yeast begin to grow

on C-2 compounds like acetate (reviewed in Krukeberg and Dickinson, 2004). This cycle is important for the formation of four carbon metabolic intermediates that can feed into the TCA cycle (reviewed in Krukeberg and Dickinson, 2004). No studies to date have examined the expression profiles of genes associated with sugar metabolism in wine yeast fermenting Icewine juice and correlated these expression patterns to the rate of sugar utilization during fermentation.

2.4.2. Nitrogen Utilization

The metabolism of amino acids in yeast is dependent on their initial uptake from the growth medium. Paulsen *et al.* (1998) listed several genes encoding for proteins with amino acid transporting activity as part of a larger study involving the categorization of a number of putative and established transporters encoded in the yeast genome. In 1999, Regenberg *et al.* extensively investigated and subsequently described the substrate specificity of 15 identified amino acid permeases. Based on their results and those from other studies, some transporters are general, such as Gap1p and Agp1p, since they display broad substrate specificity by transporting all amino acids and other nitrogenous compounds across the yeast membrane. Some transporters are class-specific and exhibit narrower substrate specificity, for example, Bap2p and Bap3p, both of which are specific for transporting branched chain amino acids. Although class-specific amino acid transporters have a higher affinity for their preferred amino acid substrates than the general carriers, regulation of the expression of genes encoding for all amino acid transporters is dependent in part on available extracellular nitrogen sources (Regenberg *et al.*, 1999).

The availability and composition of nitrogen-containing compounds in grape juice can affect yeast metabolism during fermentation and ultimately the sensory characteristics of the final wine, by influencing the production of a number of aroma and spoilage compounds (Monteiro and Bisson, 1991a; reviewed in Boulton *et al.*, 1998). Grape juice contains a variety of nitrogenous compounds, which is dependent on grape variety and time of harvest (Kluba *et al.*, 1978). Ammonium ions and amino acids are the two main sources of yeast-assimilable nitrogen (YAN) compounds in grape juice, with ammonium ions making up a large percentage (up to 40%) of the total YAN. Proline and arginine are the most common nitrogenous compounds present in juice (35-60% of total amino acid content) (Beltran *et al.*, 2004; reviewed in Boulton *et al.*, 1998). However, yeast are not able to utilize proline during fermentation and so, it is important to exclude proline when determining YAN in juice. The other major nitrogen species found in unfermented grape juice are alanine, γ -aminobutyrate (GABA), glutamate, glutamine, serine and threonine (reviewed in Boulton *et al.*, 1998). High concentrations of GABA present in the juice may possibly be formed in grape postharvest (reviewed in Boulton *et al.*, 1998). GABA is among the major nitrogen sources present in grape juice that can be used by yeast as a nitrogen source during fermentation of grape juice (Monteiro and Bisson, 1991a). GABA is a non-protein amino acid, which plays a number of important roles in eukaryotic systems, including oxidative stress tolerance in *S. cerevisiae* (reviewed in Shelp *et al.*, 1999; Coleman *et al.*, 2001). Previous studies have shown that GABA may be used as a compatible osmolyte in plants subjected to salt stress (reviewed in Shelp *et al.*, 1999). Glutamate can be decarboxylated by a glutamate decarboxylase to synthesize GABA and in turn, GABA can be degraded to form succinate in a two step

reaction. No studies to date have reported on the utilization of various nitrogen sources like GABA during Icewine fermentation and how this may relate to reduced rates of fermentation and biomass accumulation.

S. cerevisiae is able to use different nitrogen sources for growth but shows selectivity towards preferred nitrogen sources like ammonia and glutamine by repressing the genes encoding for enzymes required for the utilization of poorer nitrogen sources like ornithine, GABA, allantoin and urea. A mechanism, referred to as nitrogen catabolite repression (NCR), is invoked (reviewed in Magasanik and Kaiser, 2002). Amino acids have three distinct fates in *S. cerevisiae*: direct incorporation within biosynthetic pathways, conversion to a related compound for subsequent biosynthesis, or degradation, releasing nitrogen either as a free ammonium ion or as bound nitrogen via a transamination reaction (reviewed in Boulton *et al.*, 1998). Past studies have established that *S. cerevisiae* shows selectivity towards the uptake and utilization of different nitrogen sources (Monteiro and Bisson, 1991a and 1991b). Compounds considered as good sources of nitrogen are the ammonium ion, glutamate and glutamine, all of which are specifically depleted from the medium first before poorer sources of nitrogen, like arginine, asparagine, aspartate, urea and GABA, are taken up (reviewed in Cooper, 1982; Boulton *et al.*, 1998). Despite high concentrations of nitrogen compounds in the starting juice, yeast fermenting Icewine juice take up approximately half of the available amino acids (Kontkanen *et al.*, 2004; Pigeau and Inglis, 2005). Therefore, it is unclear if differences exist in the usage of individual amino acids by yeast during an Icewine fermentation compared to a table wine fermentation.

Previous studies by Pigeau and Inglis (2005 and 2007) showed that there were clear differences the rates of sugar consumption, nitrogen utilization, glycerol and acetic acid production by wine yeast during the first week of Icewine juice fermentation relative to a diluted Icewine juice fermentation. However, there have been no studies published to date that have compared the transcriptomic differences associated with wine yeast during Icewine fermentation compared to a table wine fermentation. To understand how and why wine yeast respond differently during Icewine fermentation, the focus of my project was to identify and compare these differences in *S. cerevisiae* using cDNA microarray technology, during the first five days of fermentation. Differences in gene expression patterns on a genomic level were correlated to yeast cell growth, sugar consumption, nitrogen usage and metabolites produced by the yeast under the two fermentation conditions.

3. MATERIALS AND METHODS

3.1. YEAST STRAINS

The commercial wine yeast strain used for fermentations, *Saccharomyces cerevisiae* K1-V1116, was provided by Lallemand Inc. (Montréal, QC, Canada). Yeast strains used for the validation of DNA probe specificities for Northern analysis were deletion mutants ($\Delta ACS1$, $\Delta ACS2$ and $\Delta STL1$) of the parental strains of *S. cerevisiae* S228C BY4742 (MAT α his3 Δ 1 leu2 Δ 0 lys2 Δ 0 ura3 Δ 0) and BY4743 (MAT α his3 Δ 1 leu2 Δ 0 lys2 Δ 0 met15 Δ 0 ura3 Δ 0) obtained from Open Biosystems (Huntsville, AL, USA).

3.2. MEDIA

To validate the specificity of DNA probes prior to Northern analysis, parental and deletion mutant yeast strains were grown under inducing and non-inducing conditions specific for the expression of the gene under investigation. All deletion mutants were grown with 200 $\mu\text{g mL}^{-1}$ Geneticin (G418) (Sigma; Oakville, ON, Canada) For *STL1* probe validation, yeast cells were grown aerobically overnight in YPD medium (1% yeast extract, 2% peptone, 2% dextrose) at 30°C with shaking at 150 rpm. Osmotic stress was induced in the cells following the addition 5 M sodium chloride to a final concentration of 0.7 M NaCl and using a 30 minute exposure time prior to RNA isolation as described by Rep *et al.* (2000). For *ACS1* probe validation, yeast cells were grown aerobically for three days in YPD medium at 30°C with shaking at 150 rpm prior to RNA isolation. For *ACS2* probe validation, yeast cells were grown anaerobically at 30°C for two days in a modified YPD medium containing 4% dextrose.

3.3. ICEWINE JUICE PREPARATION AND COMPOSITIONAL ANALYSIS

Vidal Icewine juice was kindly provided by Inniskillin Wines (Niagara-on-the-Lake, ON, Canada). The juice was filtered through a series of coarse, medium and fine pore-size pads using a Bueno Vino Mini Jet filter system (Vineco; St. Catharines, ON, Canada). The juice was then sterile-filtered through a 0.22 µm membrane cartridge filter (Millipore; Etobicoke, ON, Canada) into sterile 1L bottles. The sterile juice was stored at -40°C prior to the fermentation experiments. Two fermentation conditions were used to investigate the osmotic stress response of wine yeast using this juice. One condition required that the Icewine juice remained undiluted at 40.1°Brix prior to fermentation and the other condition required the dilution of the Icewine juice to 20.2°Brix using sterile distilled water (RiOs-16; Millipore).

The initial Icewine juice was assessed for soluble solids, titratable acidity (T.A.), pH and total nitrogen content prior to inoculation and fermentation (Table 3.1). All analyses were performed in duplicate. Soluble solids of the starting unfermented juices were determined with an ABBE bench top refractometer (American Optical). The titratable acidity of the initial juices was determined using the method outlined by Zoecklein *et al.* (1996). The pH of the initial juices was measured using a Corning pH meter (model 445) first calibrated to pH 4.0, pH 7.0 and pH 10.0. Yeast assimilable amino acid nitrogen (YAN) levels were determined using the *O*-phthaldialdehyde (NOPA) assay described by Dukes and Butzke (1998). Ammonia nitrogen in the initial juice was determined in duplicate using the ammonia enzyme assay kit from Megazyme (Bray Co., Wicklow, Ireland).

Table 3.1: Icewine juice parameters.

| Parameter | Value* |
|---------------------------------|---|
| Soluble solids | 40.2°Brix |
| Reducing sugars | 447.7 ± 8.5 g L ⁻¹ |
| Glucose | 193.0 ± 4.6 g L ⁻¹ |
| Fructose | 254.7 ± 4.1 g L ⁻¹ |
| pH | 3.6 ± 0.2 |
| Titrateable acidity (TA) | 6.8 ± 0.2 g L ⁻¹ Tartaric acid |
| Assimilable amino acid nitrogen | 388.1 ± 5.5 mg N L ⁻¹ |
| Ammonia nitrogen | 38.6 ± 3.2 mg N L ⁻¹ |
| Glycerol | 1.9 ± 0.2 g L ⁻¹ |
| Acetic Acid | 0.03 ± 0.00 g L ⁻¹ |
| Succinate | 0.05 ± 0.01 g L ⁻¹ |

*Represents mean values ± standard deviation of three independent replicate Icewine juice samples with tests performed in duplicate (n=6).

3.4. STARTER CULTURE PREPARATION AND INOCULATION

Starter cultures were prepared with the commercial yeast *Saccharomyces cerevisiae* K1-V1116 by using step-wise acclimation method. In a sterile 250 mL Erlenmeyer flask, 5.0 g of dehydrated yeast was rehydrated in 50 mL of sterile, 40°C distilled water for 15 min with gentle swirling every 5 min to encourage aeration. To acclimatize the yeast to the growth medium, 50 mL of sterile diluted Icewine juice diluted to 20°Brix was added aseptically to the rehydrated yeast and the starter culture was incubated at 25°C for 1 h in a water bath, swirling every 30 min. An additional 50 mL of undiluted Icewine juice was introduced to the starter culture achieving a final juice concentration of approximately 20°Brix. The starter culture was incubated at 20°C for 2 h and the culture was aerated by periodically swirling the flask every 30 min. A sample of the starter culture was examined for actively budding yeast cells under 40× magnification using a light microscope prior to inoculation. In sterile 0.5 L fermentation vessels fitted with air locks, 7.5 mL of the starter culture was used to inoculate 0.5 L each of Icewine juice (40°Brix) and diluted Icewine juice (20°Brix), reaching a final yeast inoculation rate of 0.5 g (dry weight) L⁻¹.

3.5. FERMENTATION AND METABOLITE ANALYSIS

Fermentations were carried out at 17°C and were monitored daily for the first week and then every other day thereafter for sugar consumption, biomass production and cell population. Fermentations were allowed to continue until approximately 200 g L⁻¹ of sugar was consumed by the yeast. All fermentations were performed in triplicate. Yeast biomass accumulation was determined using a filter retention assay. A 5 mL sample from each fermentation condition was removed and passed through a sterile, preweighed 0.45 µm cellulose nitrate membrane filter under a vacuum. The undiluted filtrate was sterile-filtered through a 0.22µm syringe filter before being stored at -30°C until metabolite analysis was performed. Filters were washed twice with deionized water and dried at 60°C for 2 days. Filters were then reweighed and biomass was determined as the difference in mass. Total cell concentrations were determined in duplicate using a haemocytometer under 40× magnification with a light microscope. Fermentation samples were diluted with 0.1 M potassium phosphate buffer (pH 7.0) prior to counting. Reducing sugar (glucose and fructose), glycerol, acetaldehyde and acetic acid levels, in samples taken from both the starting and the fermenting juices, were determined in duplicate using enzyme assay kits from Megazyme. Succinate concentrations in the finished wines were also determined in duplicate using an enzyme kit from Megazyme. Changes in yeast assimilable amino acid nitrogen levels were followed throughout the fermentations using the NOPA assay and measurements were performed in duplicate. Ethanol levels in the final wines were determined in triplicate using a Hewlett-Packard 5890 gas chromatograph with a 30 m × 0.32 mm (5%) phenyl methyl silicone column. All samples were diluted 10-fold prior to analysis and contained 1% (v/v) n-butanol as an internal

standard. XL Stat v7.1 (Addinsoft; NY., USA) was used for data analysis. Significant differences in metabolite levels were determined using a two-tailed paired t-test.

3.6. RNA EXTRACTION

The RNA extraction protocol used in this study was adapted from the protocol used by Pigeau and Inglis (2005). Juice sample volumes up to 50 mL were removed directly from the fermentation vessels and to the samples 0.01 mg mL^{-1} cyclohexamide was added prior to centrifugation. The cells were pelleted at 3800 g for 5 min at 4°C using a Sorvall RC 5C plus centrifuge (Sorvall; Newtown, CT, USA). The cells were resuspended in 10 mL of DEPC-treated distilled water and were pelleted as before. On ice, cells were resuspended in $375 \text{ }\mu\text{L}$ of extraction buffer (0.1 M NaCl , 10 mM Tris-Cl ($\text{pH } 8.0$), 1 mM EDTA ($\text{pH } 8.0$) and 5% Triton X-100) and $250 \text{ }\mu\text{L}$ phenol/chloroform/isoamyl alcohol (PCI). Following the addition of 300 mg of glass beads, the suspension was vortexed at 2500 rpm for 6 min to lyse the cells. After vortexing, $6 \text{ }\mu\text{L}$ of 20% (w/v) SDS was added and the mixture was left to stand on ice for 1 h. The mixture was centrifuged at 16000 g in a Sorvall RMC-14 centrifuge for 20 min at 4°C . To precipitate the RNA, $15 \text{ }\mu\text{L}$ of 5 M NaCl and $1250 \text{ }\mu\text{L}$ absolute ethanol was added to the supernatant. The solution was allowed to sit at -30°C for 2 h prior to centrifugation at 16000 g for 20 min at 4°C . The resulting RNA pellet was resuspended in $50 \text{ }\mu\text{L}$ of DEPC-treated distilled water and stored at -80°C until use. RNA concentration was determined spectrophotometrically at 260 nm . RNA extracted from yeast cells fermenting the Icewine and diluted Icewine juice was subsequently used for microarray and Northern analysis.

3.7. MICROARRAY ANALYSIS

3.7.1. Reverse transcription and hybridization

Total RNA (30 ug) isolated from yeast was purified using the RNeasy Minelute Cleanup Kit (Qiagen; Mississauga, ON, Canada). An aliquot of purified RNA was sent to the Microarray Centre of the University Health Network in Toronto, ON, Canada (<http://www.microarrays.ca/>) to assess RNA quality and purity using an Agilent 2100 Bioanalyzer. RNA samples not showing degradation were used to generate fluorescently labeled cDNA for subsequent microarray analysis. Yeast 6.4Kv6 cDNA microarray slides were purchased from The Microarray Center. The slides contain 6240 characterized and unknown yeast ORFs spotted in duplicate, together with positive and negative control spots. cDNA copies of the RNA transcripts obtained from yeast fermenting dilute juice or Icewine juice were generated using a modified direct labeling protocol provided by The Microarray Center. One microarray hybridization experiment was performed per fermentation replicate. Cyanine dyes were obtained from Perkin-Elmer (Boston, MA, USA). Fluorescently-labeled first-strand cDNA probes were synthesized from 20 µg of total purified RNA in a 40 µL reaction mix containing: 8.0 µL of 5X first strand buffer (Invitrogen; Burlington, ON, Canada), 2.0 µL 100 pmole µL⁻¹ AncT primer (5' T20VN 3'), 3.0 µL of 6.67 mM each of dATP, dGTP and dTTP; 1.0 µL 2 mM dCTP, 4.0 µL 0.1 M DTT, 5.0 µL of 5 ng artificial Arabidopsis transcript, 1.0 µL 1 mM cyanine 3-dCTP (Cy3, diluted Icewine juice samples) or cyanine 5-dCTP (Cy5, Icewine juice samples). The reaction mixture was first incubated at 65°C for 5 min and then cooled to 55°C for an additional 5 min to denature the RNA and anneal the AncT primer. Reverse transcription was initiated following the addition of 2 µL of Superscript III reverse transcriptase

(Invitrogen) and the reactions were incubated at 55°C for 2 h. The reaction was stopped by adding 4 µL of 50 mM EDTA (pH 8.0) and the RNA templates were hydrolyzed by adding 2 µL of 10 N NaOH to the reaction, followed by an incubation at 65°C for 20 min. The reaction was then neutralized by adding 4 µL of 5 M acetic acid. The labeled cDNA probes were pooled and purified using the CyScribe GFX Purification Kit (Amersham Biosciences; Piscataway, NJ, USA) and concentrated to a final volume of 5 µL by evaporation under vacuum using a Savant DNA110 Speed Vac. The concentrated probe was added to 80 µL of rehydrated DIG Easy Hyb (Roche) containing 5 µL of calf thymus tRNA and 5 µL yeast tRNA. The solution was heated to 65°C for 2 min and then allowed to fully cool to room temperature. The solution was applied to a yeast microarray slide covered with a Lifterslip (ErieScientific; Portsmouth, NH, USA). The slides were placed in a hybridization chamber overnight at 37°C. Following hybridization, the slides were briefly dipped in a 1× SSC solution to remove the Lifterslips. The slides were then washed three times in 1× SSC solution containing 0.1 % (w/v) SDS in 15 min intervals at 50°C with occasional agitation. After the washes were complete, the slides were briefly rinsed in 1× SSC followed by a second rinse in 0.1× SSC. The slides were dried by centrifugation at 40 g in a Sorvall RC 5C plus centrifuge for 10 min.

3.7.2. Image acquisition and data analysis

The slides were scanned using a PerkinElmer ScanArray Lite confocal laser scanner with excitation/emission wavelengths of 550 nm/570 nm for Cy3 and 649 nm/670 nm for Cy5. A preliminary scan was performed at a resolution of 50 µm to adjust laser power and photomultiplier tube (PMT) settings using the Line Scan tool until the fluorescent signal intensity of the Arabidopsis control spots was equal in each dye

channel. A final scan was performed at a resolution of 10 μm . For each dye channel, the fluorescent gene spot signal intensities were stored as separate 16-bit grayscale TIFF image files and were quantified with ScanArray Express software v3.0 (PerkinElmer) using the adaptive spot finding method. The background-adjusted signal intensities were calculated by subtracting the median local background intensity from the median signal intensity per gene spot. The adjusted signal intensities were then normalized by the Lowess (locally weighted linear regression) method. Gene spots having a normalized signal-to-background value less than 500 units were flagged and removed from subsequent analysis. Poor quality spots were flagged manually by the user and were also removed from the dataset. Of those gene spots passing the quality criteria, normalized gene expression values were reported as \log_2 transformed median Cy3 and Cy5 signal intensity values and exported as a tab delimited text data file in Microsoft Excel 2000 (Microsoft Corporation; Redmond, WA, USA). The text files were imported into GeneMaths XT v.1.5 (Applied Maths; Austin, TX, USA), which is a microarray statistical and datamining software package. A paired t-test was performed to determine significant differences in median Cy3 and Cy5 signal intensity values for each gene spot. From these values, the Cy5/Cy3 signal intensity ratios were calculated to determine fold differences in gene expression change between fermentation conditions. Gene spots passing the paired t-test ($P \leq 0.05$) and showing at least a two-fold change in the level of expression between fermentation conditions were subjected to datamining. GeneMaths can link directly to the *Saccharomyces* Genome Database (SGD) (<http://www.yeastgenome.org/>). Additional information using the Gene Ontology (GO) Slim term annotations for each gene in the yeast genome catalogued by the SGD was

retrieved and assigned to the appropriate genes within data sets (Ashburner *et al.*, 2000; Dwight *et al.*, 2002). These annotations provided information on the product of a given gene, such as its biological process, molecular function, cellular component and biochemical pathway association within the yeast cell.

3.8. VALIDATION OF MICROARRAY RESULTS USING NORTHERN ANALYSIS

3.8.1. DNA Probe Design

DNA probes for *STL1*, *ACS1* and *ACS2* were designed by the author against regions of low homology to related genes and gene sequences were obtained using the BLAST analysis search tool from the *Saccharomyces* Genome Database (SGD). Forward and reverse primers for each gene were designed to amplify these regions using PCR. PCR was carried out in a PTC-200 DNAEngine thermocycler (MJ Research; Waltham, MA, USA) using the following parameters: 30 cycles of 94°C for 40 s, 55°C for 60 s and 72°C for 90 s. The probe sequences are summarized in Table 3.2. The *STL1* primers (forward and reverse) amplified a 550 bp fragment, the *ACS1* primers amplified a 300 bp fragment and the *ACS2* primers amplified a 250 bp fragment. The DNA probe for *ALD3* was previously designed by Pigeau and Inglis in 2005. The primers for *ALD3* amplified a 259 bp fragment. The forward *ITS1* primer and the reverse *ITS4* primer, designed by Esteve-Zarzoso *et al.* (1999), amplified a DNA fragment size of 850 bp. The PCR products were run on a 1% (w/v) agarose gel to verify probe size and were visualized under UV light using a BioRad Gel-doc 1000 system and accompanying software. Probes were purified with a Qiagen QIAquick gel extraction kit.

Table 3.2: Primer sequences used for Northern analysis probe design

| Probe | Sequence |
|-------------|---|
| <i>STL1</i> | FWD 5'-TCAAAGGCAAATTTATAAGCAGAAC-3' REV 5'-CCAAAATCAATCCAATAAGCAATCA-3' |
| <i>ACS1</i> | FWD 5'-CAAAACTAGAAGAACAGTCAAGTGA-3' REV 5'-CTGCCCCGTTTTAGGGTCTGGGATGA-3' |
| <i>ACS2</i> | FWD 5'-TTTATGAAGCTCACAACGTAAAGGC-3' REV 5'-ATGTCTGTCAACACAATTGTATGAT-3' |
| <i>ALD3</i> | FWD 5'TGGTTATGGTTCCGTTGTGGGGAAA-3' REV 5'-AAACTCTTGAGTTTGCAGTGCAGAT-3' |
| <i>ITS1</i> | FWD 5'-TCCGTAGGTGAACCTGCGG -3' |
| <i>ITS4</i> | REV 5'-TCCTCCGCTTATTGATATGC -3' |

3.8.2. Northern Analysis

Unpurified (50 µg) total RNA samples extracted from one fermentation trial were electrophoresed at 95 V for 4.5 h in a 1.25% (w/v) agarose gel containing 18% (v/v) formaldehyde and blotted onto a positively charged nylon membrane (Boehringer Mannheim; Darmstadt, Germany) using capillary transfer. RNA was cross-linked to the membrane using a Hoefer UVC 500 UV crosslinker (Hoefer, Inc.; CA, USA). Probes were labeled with [α -³²P]-dATP (Perkin Elmer) using a Boehringer Mannheim random primed labeling kit and 20 ng of labeled probe was hybridized to the bound RNA following an overnight incubation at 42°C in 50 mL of hybridization solution containing 5× SSC, 5× Denhardt's solution, 1% (v/v) SDS, 50% (v/v) formamide and 0.1 mg mL⁻¹ sheared salmon sperm DNA. The membrane was washed twice with 50 mL volumes of 2× SSC containing 0.1% (v/v) SDS for 20 min at room temperature and washed twice with 50 mL volumes of 0.2× SSC, also containing 0.1% (v/v) SDS, for 15 min at 50°C. The probed membrane was exposed to a Fujifilm phosphorimaging screen for up to seven days and the screen was scanned using a Fujifilm FLA-3000 phosphorimager with a 633

nm helium-neon laser. The subsequent images were quantified with Fujifilm Image Gauge software (v.4.0). Gene expression levels were normalized to the signal intensities derived from a rDNA region spanning the 5.8S rRNA gene and flanking internal transcribed spacers (ITS) 1 and 2.

3.9. MEASURING AMINO ACID CONCENTRATION USING HPLC

Individual primary amino acids, including γ -aminobutyrate (GABA) and imino acids (proline), in Icewine and diluted Icewine juice samples were separated and quantified using the Agilent 1100 Series HPLC following their conversion to their respective *O*-phthaldialdehyde (OPA) and 9-fluorenylmethyl chloroformate (FMOC) derivatives. Juice samples were diluted with water prior to preparing the derivatization reactions using the Agilent 1313A autosampler according to the manufacturer's instructions. 0.5 μ L of each derivatized sample was injected into a 4.6×150 mm, 3.5 μ m ZORBAX Eclipse-AAA column. The OPA- and FMOC-derivatized amino acids were detected at 340 nm using a G1315A Fluorescence Detector (FLD). The concentration of each amino acid present per juice sample was determined in singlet for each of the three fermentation replicates and expressed in mg N L⁻¹.

4. RESULTS AND DISCUSSION

4.1. MICROARRAY ANALYSIS OVERVIEW

4.1.1. Results - Microarray

The genomic expression profiles of the commercial wine yeast *Saccharomyces cerevisiae* K1-V1116 fermenting Icewine juice were compared to those generated by yeast cells fermenting diluted Icewine juice using cDNA microarray analysis. All fermentations were carried out in triplicate and one microarray experiment was performed for each of the days examined per fermentation replicate. The expression of approximately 6240 open reading frames (ORFs) in yeast was compared between juice conditions during the first five days of fermentation which corresponded to the start of the early exponential growth phase until cells entered to early stationary phase. Gene expression was considered to be significantly up- or downregulated in the Icewine juice condition if the difference in gene expression passed a paired t-test ($P \leq 0.05$) and showed at least a two-fold change in the level of expression between fermentation conditions. A small fraction of the number of genes within the yeast genome displayed significant changes in expression in response to the fermentation of Icewine juice (Table 4.1.1). Of these genes, the vast majority showed a difference in expression between juice conditions resulting in an up- or downregulation of less than five-fold in the Icewine juice fermentation. The total number of up- and downregulated genes was approximately equivalent on day two of the fermentations. The total number of genes downregulated was slightly higher than those upregulated for each of the remaining fermentation days analyzed (Table 4.1.1). The total number of genes differentially expressed in yeast

Table 4.1.1: Distribution of the differentially expressed genes in *S. cerevisiae* fermenting Icewine juice and dilute Icewine juice.

| Expression | Fold Change* | Number of Genes | | | |
|---------------|---------------------|-----------------|-------|-------|-------|
| | | Day 2 | Day 3 | Day 4 | Day 5 |
| Upregulated | 2 to <5 | 120 | 80 | 115 | 37 |
| | ≥5 to <10 | 9 | 13 | 21 | 11 |
| | ≥10 | 1 | 6 | 12 | 5 |
| | Total upregulated | 130 | 99 | 148 | 53 |
| Downregulated | 2 to <5 | 120 | 131 | 154 | 55 |
| | ≥5 to <10 | 19 | 12 | 24 | 9 |
| | ≥10 | 6 | 1 | 3 | 8 |
| | Total downregulated | 145 | 144 | 181 | 72 |
| Total | | 275 | 243 | 329 | 125 |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

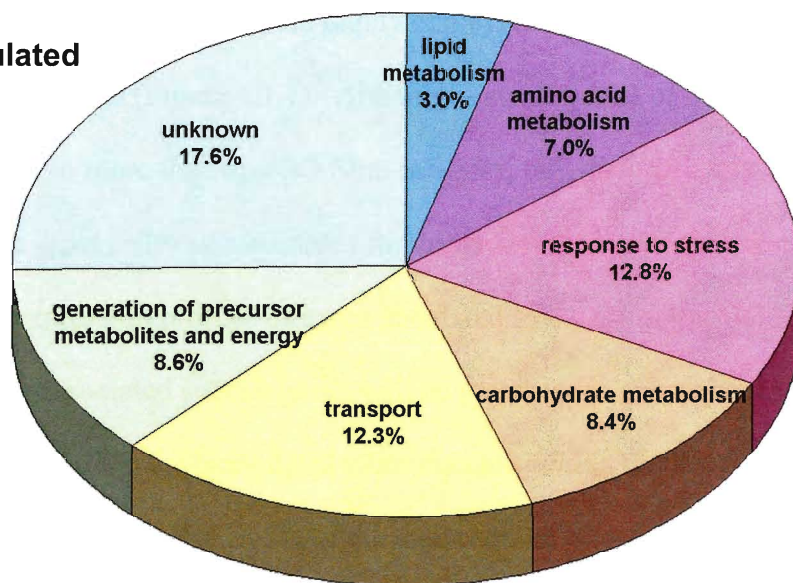
between fermentation conditions reached a maximum on day four (329 genes) which represents about 5% of the genome. The total number began to decline on day five (125 genes) as cells entered stationary phase (section 4.2.1.1.).

Genes were also characterized according to Gene Ontology (GO) Slim terms of the *Saccharomyces* genome database (SGD). Those genes identified as being differentially expressed at least two-fold between fermentation conditions were grouped together into broad categories based on common biological gene function within the yeast cell. Table 4.1.2 lists the distribution of the genes differentially expressed in yeast fermenting Icewine juice based on their 2007 GO Slim biological function annotations. The number of gene expression changes was transient and occurred at a low frequency throughout the early stages of the Icewine fermentation for genes associated with GO Slim categories related to cell growth and reproduction. This category included cellular functions related to budding, cytokinesis, morphogenesis and sporulation (Table 4.1.2). In contrast, a greater number of yeast genes whose expression was significantly affected by fermentation of Icewine juice and consistently displayed overall up- or downregulation throughout the first five days of fermentation could be divided into seven main biological categories as illustrated in Figure 4.1.1. When the number of genes differentially expressed more than two-fold between fermentation conditions was divided by the total number of genes either up- or downregulated from days two to five of the fermentations in each GO Slim category, it was found that the overall expression of genes associated with the generation of precursor metabolites and energy, lipid metabolism, amino acid metabolism, carbohydrate metabolism and transport were expressed to a lesser degree in yeast fermenting Icewine juice relative to cells fermenting the dilute

Table 4.1.2: Distribution of GO Slim terms for differentially expressed genes in *S. cerevisiae* fermenting Icewine juice.

| GO Slim Term | Day 2 | | Day 3 | | Day 4 | | Day 5 | |
|--|-------|------|-------|------|-------|------|-------|------|
| | Up | Down | Up | Down | Up | Down | Up | Down |
| amino acid and derivative metabolism | 8 | 17 | 10 | 10 | 8 | 14 | 4 | 6 |
| carbohydrate metabolism | 14 | 11 | 8 | 24 | 7 | 30 | 7 | 9 |
| cell budding | 0 | 1 | 0 | 2 | 0 | 2 | 0 | 2 |
| cell cycle | 6 | 6 | 4 | 7 | 9 | 8 | 2 | 3 |
| cell homeostasis | 1 | 1 | 2 | 6 | 2 | 6 | 1 | 6 |
| cell wall organization and biogenesis | 3 | 3 | 3 | 7 | 6 | 9 | 1 | 6 |
| cellular respiration | 6 | 2 | 0 | 17 | 0 | 25 | 1 | 2 |
| conjugation | 1 | 2 | 1 | 4 | 1 | 4 | 0 | 0 |
| cytokinesis | 0 | 1 | 0 | 1 | 1 | 1 | 1 | 2 |
| cytoskeleton organization and biogenesis | 0 | 1 | 0 | 2 | 1 | 3 | 1 | 3 |
| DNA metabolism | 20 | 4 | 4 | 8 | 9 | 11 | 2 | 2 |
| electron transport | 2 | 1 | 0 | 13 | 0 | 12 | 0 | 1 |
| generation of precursor metabolites and energy | 19 | 16 | 6 | 49 | 5 | 53 | 7 | 17 |
| lipid metabolism | 6 | 21 | 4 | 2 | 3 | 11 | 0 | 1 |
| meiosis | 2 | 5 | 3 | 5 | 6 | 5 | 2 | 0 |
| membrane organization and biogenesis | 3 | 2 | 0 | 1 | 1 | 2 | 0 | 0 |
| morphogenesis | 2 | 2 | 1 | 4 | 3 | 4 | 0 | 5 |
| nuclear organization and biogenesis | 0 | 2 | 0 | 0 | 0 | 0 | 0 | 0 |
| organelle organization and biogenesis | 11 | 9 | 9 | 12 | 25 | 18 | 3 | 7 |
| other | 7 | 11 | 6 | 6 | 12 | 11 | 8 | 4 |
| protein biosynthesis | 20 | 9 | 11 | 6 | 5 | 11 | 2 | 7 |
| protein catabolism | 2 | 3 | 1 | 1 | 1 | 2 | 1 | 3 |
| protein modification | 7 | 8 | 4 | 8 | 10 | 11 | 1 | 2 |
| pseudohyphal growth | 1 | 0 | 1 | 0 | 2 | 1 | 0 | 1 |
| response to stress | 10 | 12 | 15 | 12 | 18 | 14 | 12 | 12 |
| ribosome biogenesis and assembly | 0 | 1 | 4 | 1 | 16 | 1 | 2 | 2 |
| RNA metabolism | 3 | 5 | 5 | 5 | 15 | 5 | 3 | 5 |
| signal transduction | 0 | 6 | 2 | 1 | 2 | 3 | 2 | 0 |
| sporulation | 0 | 5 | 2 | 3 | 2 | 3 | 2 | 3 |
| transcription | 6 | 2 | 5 | 6 | 14 | 7 | 4 | 2 |
| transport | 18 | 28 | 13 | 25 | 17 | 31 | 5 | 13 |
| unknown | 16 | 19 | 22 | 23 | 32 | 29 | 6 | 7 |
| vesicle-mediated transport | 2 | 5 | 3 | 3 | 5 | 7 | 2 | 3 |
| vitamin metabolism | 7 | 7 | 2 | 2 | 5 | 4 | 2 | 2 |

(A) Upregulated



(B) Downregulated

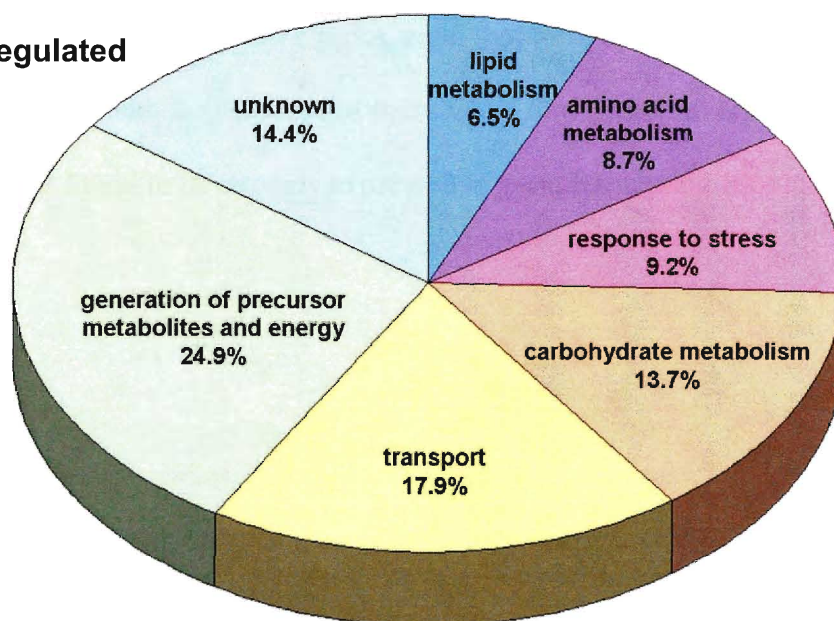


Figure 4.1.1: The major Gene Ontology (GO) Slim term categories containing genes differentially expressed in *S. cerevisiae* fermenting Icewine juice. The overall percentage of genes associated with each of the GO Slim categories containing genes that were found to be (A) upregulated or (B) downregulated in yeast fermenting Icewine juice.

juice. In contrast, more genes involved with the response to stress and those with unknown function were found to be more highly expressed on days two to five of the Icewine juice fermentation (Figure 4.1.1). Although a small subset of these genes may have been annotated to more than one GO Slim category, only their primary biological function within the yeast cell was considered for this calculation. A more detailed description of the expression profiles of genes annotated to the six major GO Slim categories and their associated subcategories will be presented in subsequent sections. Very few genes were up- or downregulated more than ten-fold in the Icewine juice condition, which represents less than 5% of the total number of genes differently expressed between fermentations (Table 4.1.1). Table 4.1.3 lists the genes with changes in expression calculated to be more than ten-fold between juice conditions on one or more days of the fermentations. Of these genes, eight that were upregulated more than ten-fold were associated with the response to stress while four genes having no known function to date were found to be strongly expressed in yeast fermenting Icewine juice.

Table 4.1.3: Genes up- or downregulated more than 10-fold in *S. cerevisiae* fermenting Icewine juice.

| ORF | Gene | Description | GO Slim Category |
|----------------------|--------|-------------------------------------|--|
| Upregulated | | | |
| YCR010C | ADY2 | acetate transporter | transport |
| YMR169C | ALD3 | aldehyde dehydrogenase | response to stress |
| YHR053C | CUP1-1 | metallothionein; copper ion binding | response to stress |
| YHR055C | CUP1-2 | metallothionein; copper ion binding | response to stress |
| YIL053W | GPP1 | glycerol-3-phosphatase | response to stress |
| YER062C | GPP2 | glycerol-3-phosphatase | response to stress |
| YOL151W | GRE2 | oxidoreductase activity | response to stress |
| YNL036W | NCE103 | carbonate dehydratase | response to stress |
| YGR159C | NSR1 | rRNA processing | RNA metabolism |
| YJR095W | SFC1 | succinate-fumarate transporter | transport |
| YMR175W | SIP18 | phospholipid binding | response to stress |
| YHR139C | SPS100 | spore wall maturation protein | sporulation |
| YDR536W | STL1 | glycerol proton symporter | transport |
| YBR281C | | unknown | unknown |
| YGL157W | | unknown | unknown |
| YHR087W | | unknown | unknown |
| YOL150C | | unknown | unknown |
| Downregulated | | | |
| YAL054C | ACS1 | acetyl-CoA synthetase | generation of metabolites and energy |
| YLR153C | ACS2 | acetyl-CoA synthetase | generation of metabolites and energy |
| YGR204W | ADE3 | formate-tetrahydrofolate ligase | other |
| YDR384C | ATO3 | ammonium transporter | transport |
| YBL101C | ECM21 | unknown | cell wall organization and biogenesis |
| YMR212C | EFR3 | unknown | morphogenesis |
| YGR254W | ENO1 | enolase | carbohydrate metabolism |
| YPL281C | ERR2 | phosphopyruvate hydratase activity | unknown |
| YOR388C | FDH1 | formate dehydrogenase | vitamin metabolism |
| YPL275W | FDH2 | formate dehydrogenase | other |
| YKR002W | PAP1 | polynucleotide adenylyltransferase | RNA metabolism |
| YGR087C | PDC6 | pyruvate decarboxylase | carbohydrate metabolism |
| YBR229C | ROT2 | alpha-glucosidase | cell wall organization and biogenesis |
| YLR410W | VIP1 | unknown | cytoskeleton organization and biogenesis |
| YIL057C | | unknown | unknown |
| YPL276W | | unknown | unknown |

4.1.2. Discussion - Microarray Overview

Past genomic profiling studies have extensively investigated the impact of salt and sorbitol-induced hyperosmotic stress on the transcriptome of laboratory strains of *S. cerevisiae* (Posas *et al.*, 2000; Rep *et al.*, 2000; Yale and Bohnert, 2001). A large scale microarray study revealed that the expression of a substantial group of nearly 900 genes is affected when yeast are challenged with a variety of environmental stresses, including hypo- and hyperosmotic stress, oxidative stress and nutrient depletion (Gasch *et al.*, 2000). Causton *et al.* (2000) found that 10% of genes within the yeast genome affected by transient stress respond in a similar manner and are expressed collectively under many different stress conditions. Genes that are induced or repressed by stress and are within this “common environmental response” or CER group are involved in carbohydrate metabolism, the generation of energy, response to reactive oxygen species, response to stress, ion homeostasis and protein synthesis. The results from the microarray data obtained in the present study revealed that 2-5% of genes within the genome of the commercial wine yeast K1-V1116 were differentially expressed between Icewine and diluted Icewine juice fermentation conditions. The bulk of these genes are involved in carbohydrate metabolism, transport and amino acid and lipid metabolism. The majority of genes that also demonstrated a difference in transcriptional response between fermentation conditions are associated with the generation of precursor metabolites and energy which were observed to be downregulated in cells fermenting Icewine juice while a greater proportion of genes associated with the response to stress, including osmotic stress, were more highly expressed relative to the table wine fermentation. In contrast to the transcriptional response of laboratory yeast strains to individual, transient

environmental stresses (Gasch *et al.*, 2000; Causton *et al.*, 2001), wine yeast cells under oenological conditions are faced with a continually changing environment and must adapt accordingly to several simultaneous stresses. It has been previously shown that solutes in Icewine juice in addition to sugar place yeast under incredible hyperosmotic stress during fermentation (Kontkanen *et al.*, 2004; Pigeau and Inglis, 2005). The transcriptional response by a wine yeast strain to stress under fermentative conditions is likely to be drastically different from that of an aerobically grown laboratory strain (reviewed in Pretorius, 2000). Only 28 out of the 186 genes induced in a laboratory yeast strain briefly exposed to salt and sorbitol hyperosmotic stress (Rep *et al.*, 2000) were found to be similarly induced in wine yeast during the first five days of Icewine juice fermentation relative to table wine fermentation. These genes were associated with the metabolism of glycerol and amino acids, maintaining redox balance and responding to cell stress. However, a number of genes induced under salt and/or sorbitol stress but not during Icewine fermentation were associated with gene functional categories related to signal transduction, proteolysis, vacuolar biogenesis in addition to trehalose and glycogen metabolism. The initial stresses of high osmotic pressure and low pH, coupled with the additional stress from ethanol as the fermentation progresses are involved in wine yeast stress response during fermentation and have been monitored at the transcriptomic level (Perez-Torrado *et al.*, 2002; Rossignol *et al.*, 2003). Gene expression profiling studies involving the adapted stress response of a wine yeast strain fermenting synthetic grape juice or the initial stress response to high sugar stress within a chaptalized grape juice environment similarly identified a large subset of genes differentially expressed under fermentative conditions that were associated with nutrient metabolism, transport and

response to stress (Rossignol *et al.*, 2003; Erasmus *et al.*, 2003). In a 2008 study by Marks *et al.*, transcriptional analysis of the stress response of a commercially available wine yeast strain throughout table wine fermentation identified a group of 223 highly expressed genes. This gene group was later designated as the fermentation stress response (FSR) and genes within this group were subsequently categorized based on their GO biological annotations. From the FSR, only 11 genes in my study induced throughout table wine fermentation were also found to be similarly expressed during Icewine fermentation. However, the majority of FSR genes expressed in the study by Marks *et al.* (2008) were associated with GO biological process categories that were not part of the majority of genes differentially expressed between Icewine and diluted juice fermentations. Of the FSR genes, the highly represented GO categories identified during table wine fermentation included genes encoding for uncharacterized proteins (28%), organelle organization and biogenesis (15%), protein modification processes (13%) and RNA metabolic processes (12%). However, FSR genes related to stress response (11%) and transport (18%) were also expressed during fermentation. In agreement with this study, genes involved with the response to stress and transport were found to be highly expressed during Icewine juice fermentation relative to the dilute juice fermentation. However, unlike the results reported in Marks *et al.* study, the majority of genes differentially expressed between Icewine and dilute juice fermentations were associated with the generation of precursor metabolites and energy, lipid metabolism, amino acid metabolism and carbohydrate metabolism.

The microarray experiments have clearly provided a wealth of information on the transcriptional differences associated with a commercial wine yeast strain (K1-V1116)

fermenting Icewine and diluted Icewine juice. Following statistical and biological gene function analysis, the resulting gene expression profile patterns were used in this study as a starting point to identify metabolic pathways related to the use of nutrients and the production of yeast metabolites that were significantly up- or downregulated during Icewine fermentation. Based on these results, a more detailed analysis of the key genes associated with these pathways will be discussed in subsequent chapters of this thesis. Transcriptional differences of genes linked to sugar and amino acid metabolism will be correlated to yeast cell growth and nutrient uptake in Sections 4.2 and 4.3. Also, the transcriptional differences in glycerol and acetic acid transport and metabolism related genes identified through microarray analysis and verified with Northern analysis will be correlated to differences in glycerol and acetic acid production in Sections 4.4 and 4.5. Section 4.6 will focus on the changes in expression of a number of stress response genes observed throughout the first five days of Icewine juice fermentation.

4.2. CARBOHYDRATE METABOLISM

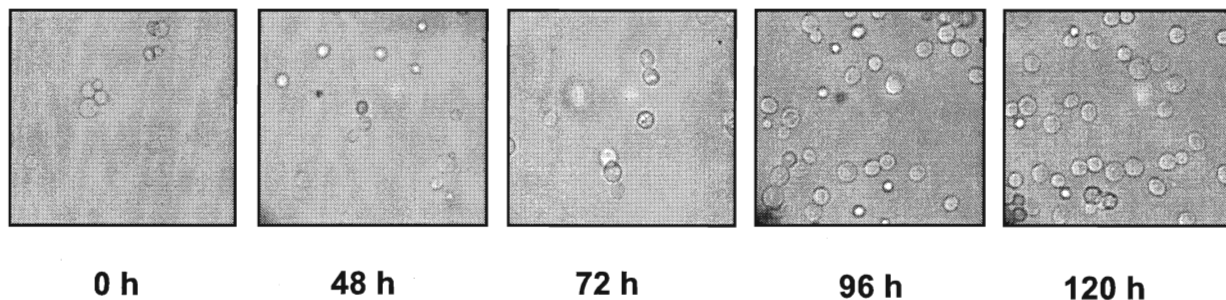
4.2.1. Results - Carbohydrate Metabolism

4.2.1.1. Yeast Cell Growth

The high level of osmotic stress placed on fermenting yeast due to the presence of concentrated solutes in Icewine juice had an overall negative impact on cell growth. In comparison to the diluted Icewine juice fermentation condition, increased extracellular osmotic pressure exerted on yeast cells fermenting Icewine juice contributed to decreased cell size and biomass accumulation, as well as lengthened the duration of the lag phase and increased the time elapsed before yeast cells began to actively bud and proliferate

(Figures 4.2.1 and 4.2.2). Icewine juice fermenting cells were visibly smaller in size due to the loss of cytosolic water and turgor pressure compared to their diluted Icewine juice fermenting counterparts up to 96 h into the fermentations (Figure 4.2.1). Up to this 48 h timepoint, cells fermenting Icewine juice were not actively budding and remained in late lag phase, while cells fermenting the dilute juice were rapidly budding and had already reached midexponential growth phase (Figure 4.2.2A). Yeast cell growth in both juice conditions reached stationary phase approximately 120 h into the fermentations. The diluted fermentation achieved a peak cell concentration of 1.1×10^8 cells mL⁻¹, whereas the high sugar fermentation accumulated a cell concentration of 6.4×10^7 cells mL⁻¹ (Figure 4.2.2A). This corresponded to a 2-fold higher peak biomass accumulation in the dilute juice fermentation in comparison to the Icewine juice fermentation, which also showed an overall slower rate of biomass accumulation (Figure 4.2.2B). These results agree with previous observations demonstrating that yeast subjected to increasing levels of osmotic stress have diminished cell volume and delayed budding (Blomberg and Adler, 1992).

(A)



(B)

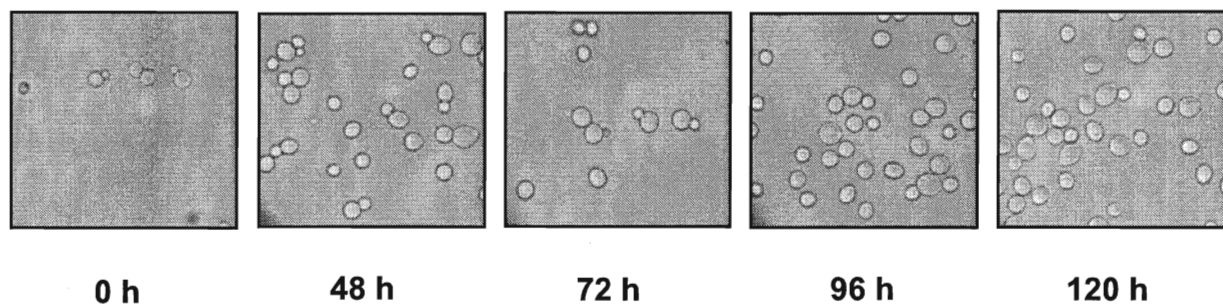


Figure 4.2.1.: Yeast cell morphology during fermentation. Yeast cells fermenting Icewine juice (A) and diluted Icewine juice (B) were visualized daily under 40× magnification.

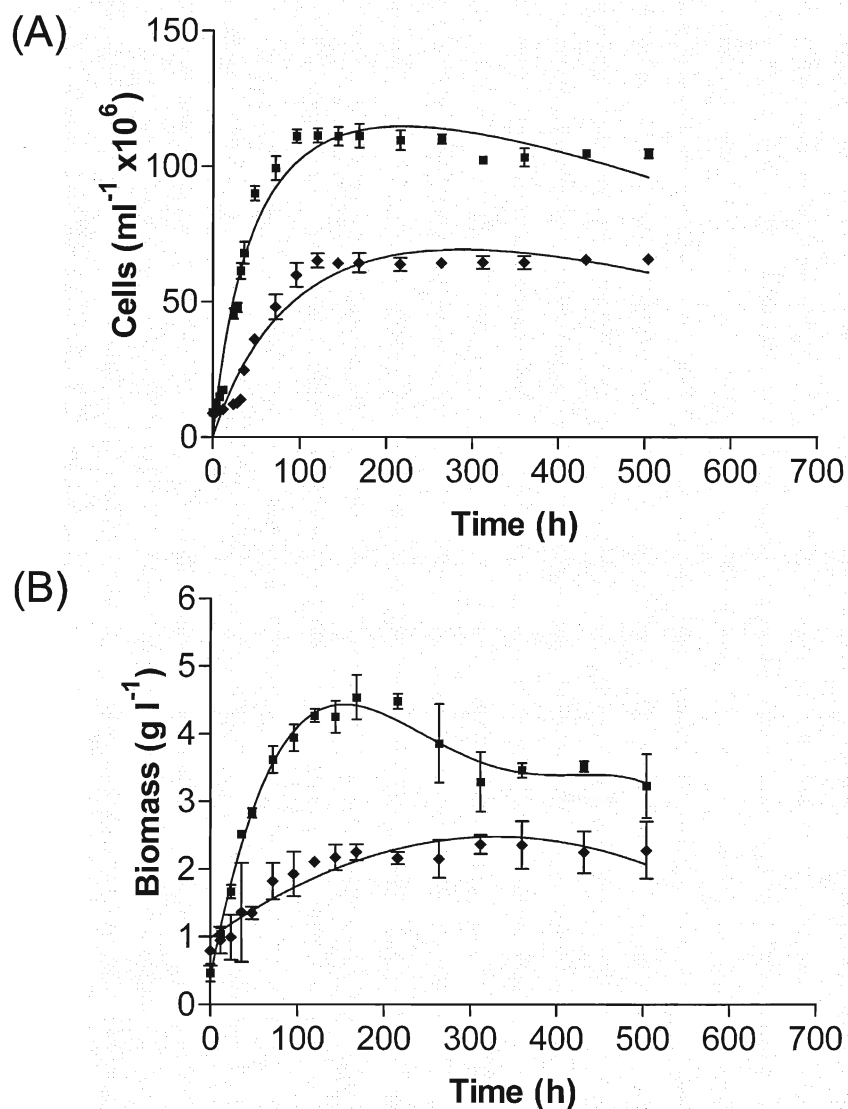


Figure 4.2.2.: Yeast cell growth and biomass accumulation. Fermentations of Icewine juice (♦) and diluted Icewine juice (■) were monitored daily for total cell counts (A) and biomass accumulation (B). The values represent the average \pm standard deviation of the mean of triplicate fermentations.

4.2.1.2. Sugar Consumption

At the start of the fermentations, there was approximately twice the amount of total sugar present in Icewine juice when compared to the diluted juice (Figure 4.2.3A). Of the reducing sugars, the unfermented Icewine and diluted juices had a lower initial glucose concentration in comparison to the fructose concentration prior to inoculation (Figures 4.2.3B and C). At the end of the fermentations, only about half of the available fermentable sugars were consumed by Icewine juice fermenting yeast in twice the amount of time it took the dilute juice cells to consume the same amount of sugar (Figure 4.2.3A). When comparing the fermentation rate between juice conditions, the high concentration of sugar and other solutes in the Icewine juice appeared to impede the ability of yeast to ferment this juice. There was a slight lag in the rate of sugar consumption during the first 48 h of the Icewine juice fermentation. The reducing sugar concentration steadily declined throughout the duration of the Icewine juice fermentation with yeast only consuming $187.8 \pm 2.8 \text{ g L}^{-1}$ of the $447.7 \pm 8.5 \text{ g L}^{-1}$ of available sugars in 576 h. In comparison, the dilute juice fermenting cells demonstrated a considerably faster fermentation rate consuming $186.8 \text{ g} \pm 3.1 \text{ L}^{-1}$ of sugar in 216 h and ultimately consuming all available reducing sugars ($221.9 \pm 0.6 \text{ g L}^{-1}$) in 312 h (Figure 4.2.3A). Additionally, there was a difference in the amount of glucose and fructose consumed by yeast between fermentation conditions. Approximately 2.5-fold more fructose ($183.9 \pm 0.3 \text{ g L}^{-1}$) remained unfermented compared to glucose ($73.9 \pm 0.8 \text{ g L}^{-1}$) in 576 h by yeast in the Icewine juice condition (Figures 4.2.3B and C). In contrast, all of the available glucose and fructose was consumed at the end of the dilute juice fermentation. Furthermore, yeast cells in both Icewine and diluted juice displayed a clear difference in

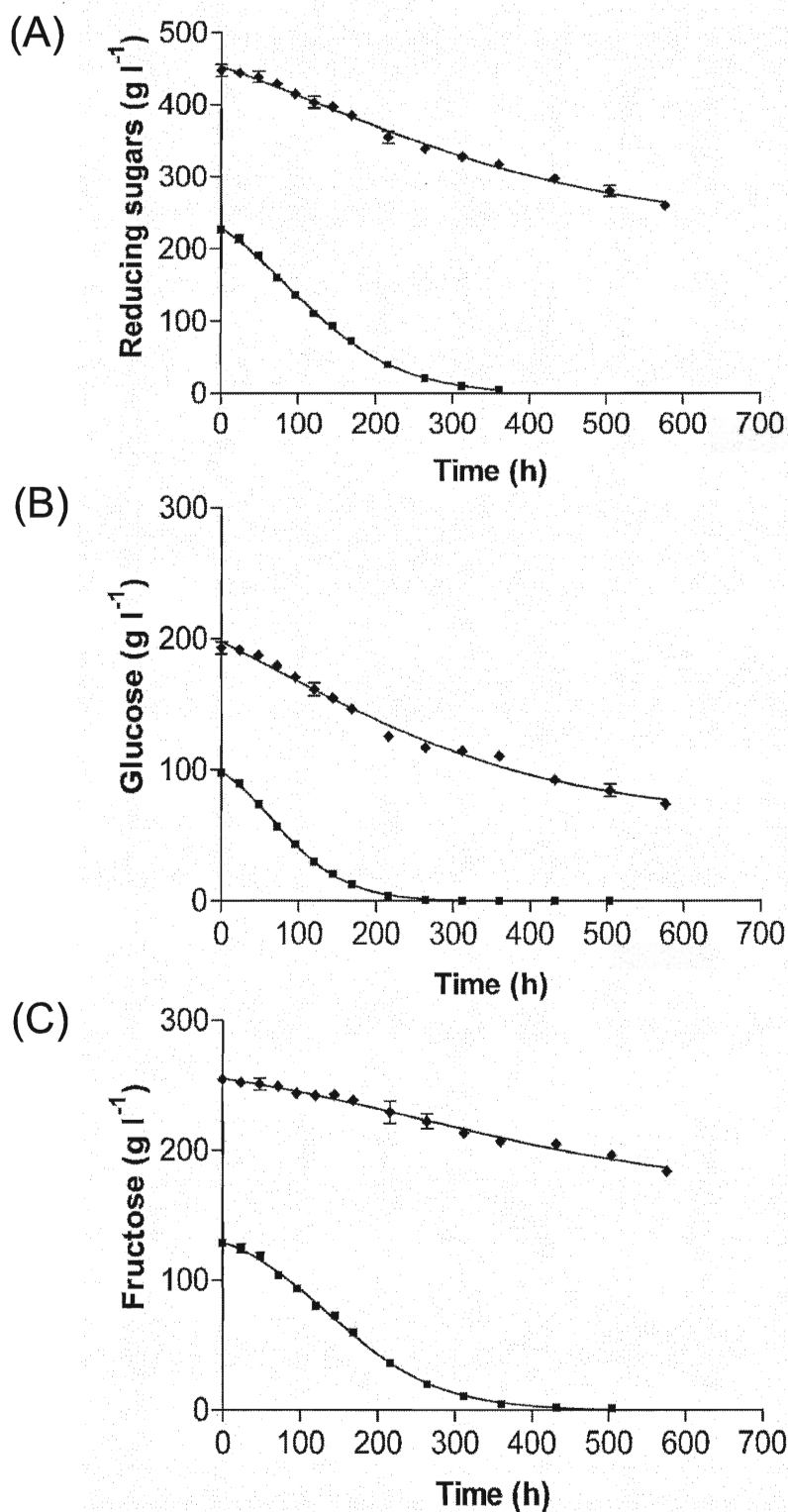


Figure 4.2.3: Yeast sugar consumption. Fermentations of Icewine juice (\blacklozenge) and diluted Icewine juice (\blacksquare) were monitored daily for total reducing sugars (A), glucose (B) and fructose (C) throughout the course of the fermentations. Fermentations were performed in triplicate with measurements performed in duplicate. The mean values \pm standard deviation are shown.

the fermentation rates of both reducing sugars. Under both fermentation conditions, yeast appeared to preferentially take up glucose since the fermentation rate of glucose was faster than that of fructose (Figures 4.2.3B and C).

4.2.1.3. Gene expression profiles for carbohydrate metabolism

Expression of the genes encoding for proteins involved in carbohydrate metabolism were highly expressed in both fermentation conditions. From the microarray results, the routes of carbohydrate metabolism in *S. cerevisiae* containing genes that were differentially expressed between fermentation conditions are illustrated in Figure 4.2.4. Upon entry into the cell, hexoses such as glucose and fructose are converted to pyruvate via glycolysis. Pyruvate is then further metabolized to either ethanol and CO₂ by alcoholic fermentation (Figure 4.2.4A) or to oxaloacetate and/or acetyl-CoA, which can enter the tricarboxylic acid cycle (TCA) (Figure 4.2.4B). Alternatively, pyruvate can enter the glyoxylate cycle (Figure 4.2.4C). All of these pathways were evaluated in fermenting yeast using microarray technology.

4.2.1.3.1. Glycolysis and alcoholic fermentation

The first step of glycolysis is the transport of sugar molecules into the yeast cell. The sugar transporter gene family encodes for the largest family of transporters in *S. cerevisiae*. In the yeast genome, there are 20 genes identified as belonging to the *HXT* family that encode for hexose transporters (Luyten *et al.*, 2002). Six *HXT* genes were observed to be differentially expressed between fermentation conditions and all were downregulated up to nine-fold throughout the Icewine juice fermentation (Table 4.4). Both *HXT8* and *HXT10* (putative hexose transporters) were expressed at the same level in

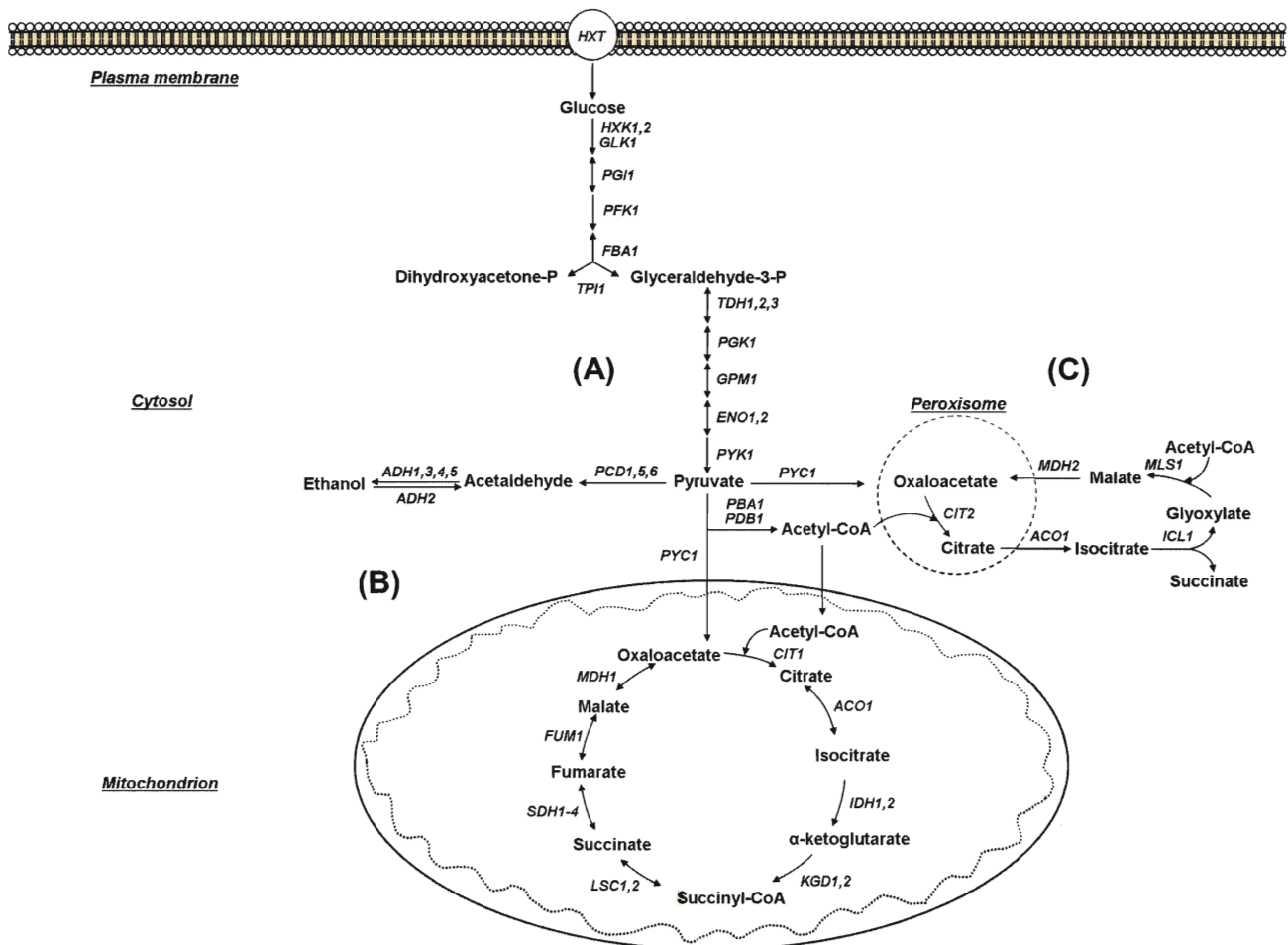


Figure 4.2.4: The pathways involved in carbohydrate metabolism differentially expressed in *S. cerevisiae* between fermentation conditions. Illustrated are the metabolic intermediates and the genes associated with (A) glycolysis and alcoholic fermentation, (B) the tricarboxylic acid (TCA) cycle and (C) the glyoxylate cycle.

both fermentation conditions on day two, along with *HXT5* (moderate affinity glucose transporter) on days three and four. The rate of sugar consumption by dilute juice fermenting cells increased considerably after day two in comparison to Icewine juice fermenting cells, which maintain a slower but steady rate of sugar consumption as seen in Figure 4.2.3A. This observation corresponded to the degree of *HXT* downregulation in Icewine juice fermenting yeast, along with a general downregulation in the genes associated with carbohydrate metabolism (Table 4.2.1).

The large majority of genes involved in the glycolytic pathway and alcoholic fermentation were differentially expressed between fermentation conditions. However, genes specifically encoding for enzymes responsible for catalyzing the energy generating, rate-limiting steps in glycolysis all showed a relatively lower fold expression in yeast under the Icewine juice condition compared to the dilute juice fermentation after day two, as recorded in Table 4.2.1. This was the case for *HXK2* (2.9-fold) encoding for hexokinase II, which catalyzes the first irreversible enzymatic step in glycolysis by transferring a phosphate group from ATP to a hexose molecule. Both hexokinase isoforms (*HXK1* and *HXK2*) were strongly downregulated in the Icewine condition relative to the dilute condition; 6.6- and 9.5-fold on day five of the fermentations, respectively. *GLK1* encodes for a glucokinase, which participates in the phosphorylation of glucose only, and it was observed to be differentially expressed in yeast between juice conditions but was expressed at a significantly higher level on day five of the Icewine juice fermentation. *PFK1* encodes for the α -subunit of phosphofructokinase, which catalyzes the second irreversible and unidirectional reaction of the glycolytic pathway, where fructose-6-phosphate is phosphorylated to form the six carbon intermediate

Table 4.2.1: Expression profile of the genes involved in carbohydrate metabolism.

| ORF | Gene | Description | Fold Change* | | | |
|---------------------------------------|-------|--|--------------|-------|-------|-------|
| | | | Day 2 | Day 3 | Day 4 | Day 5 |
| Monosaccharide Transport | | | | | | |
| YHR094C | HXT1 | low affinity glucose transporter | - | -2.4 | -4.8 | -4.6 |
| YHR092C | HXT4 | moderate affinity glucose transporter | - | -2.9 | -5.7 | -7.8 |
| YHR096C | HXT5 | moderate affinity glucose transporter | -2.2 | • | • | -4.2 |
| YJL214W | HXT8 | putative hexose transporter | • | -2.4 | -2.9 | × |
| YFL011W | HXT10 | putative hexose transporter | - | -3.9 | -9.3 | × |
| YIL171W | HXT12 | putative hexose transporter | • | -2.2 | -4.4 | × |
| Glycolysis and Alcoholic Fermentation | | | | | | |
| YFR053C | HXK1 | hexokinase I | × | - | -2.5 | -6.6 |
| YGL253W | HXK2 | hexokinase II | +2.9 | -2.2 | -2.7 | -9.5 |
| YCL040W | GLK1 | glucokinase | • | + | + | +2.4 |
| YBR196C | PGI1 | glucose-6-phosphate isomerase | • | × | × | × |
| YGR240C | PFK1 | phosphofructokinase | • | -5.7 | -5.9 | × |
| YKL060C | FBA1 | aldolase | + | -4.0 | -3.4 | • |
| YDR050C | TPI1 | triose-phosphate isomerase | + | -4.4 | -3.3 | • |
| YJL052W | TDH1 | glyceraldehyde-3-phosphate dehydrogenase | • | -6.7 | -4.7 | • |
| YJR009C | TDH2 | glyceraldehyde-3-phosphate dehydrogenase | • | -6.0 | -3.2 | -2.6 |
| YGR192C | TDH3 | glyceraldehyde-3-phosphate dehydrogenase | • | -6.7 | -5.5 | -6.6 |
| YCR012W | PGK1 | phosphoglycerate kinase | + | -7.2 | -6.7 | -6.6 |
| YKL152C | GPM1 | phosphoglycerate mutase | +5.4 | -4.1 | -3.1 | - |
| YGR254W | ENO1 | enolase | +2.8 | -12.2 | -11.6 | -13.6 |
| YHR174W | ENO2 | enolase | +2.5 | -8.6 | -7.0 | -4.0 |
| YAL038W | PYK1 | pyruvate kinase | +2.4 | -3.5 | -3.4 | • |
| YLR044C | PDC1 | pyruvate decarboxylase | +2.4 | -4.7 | -5.4 | -3.9 |
| YLR134W | PDC5 | pyruvate decarboxylase | +2.2 | -4.4 | -5.9 | • |
| YGR087C | PDC6 | pyruvate decarboxylase | +2.2 | -4.5 | -6.7 | -10.1 |
| YOL086C | ADH1 | alcohol dehydrogenase (cytosolic) | - | -4.6 | -6.5 | -3.2 |
| YMR303C | ADH2 | alcohol dehydrogenase (cytosolic) | -2.2 | -4.4 | -6.4 | -4.9 |
| YMR083W | ADH3 | alcohol dehydrogenase (mitochondrial) | -2.0 | -3.2 | • | -3.6 |
| YGL256W | ADH4 | alcohol dehydrogenase (mitochondrial) | +2.0 | × | × | × |
| YBR145W | ADH5 | alcohol dehydrogenase (cytosolic) | × | -4.3 | -4.9 | • |
| TCA Cycle | | | | | | |
| YER178W | PDA1 | pyruvate dehydrogenase | - | -2.3 | -2.4 | • |
| YBR221C | PDB1 | pyruvate dehydrogenase | • | - | - | • |
| YGL062W | PYC1 | pyruvate carboxylase | -2.3 | - | -2.4 | +2.4 |
| YNR001C | CIT1 | citrate synthase (mitochondrial) | -2.3 | - | -2.3 | • |
| YLR304C | ACO1 | aconitase | • | - | -2.0 | -7.3 |
| YNL037C | IDH1 | isocitrate dehydrogenase | + | - | - | -2.0 |
| YOR136W | IDH2 | isocitrate dehydrogenase | +2.4 | - | • | • |
| YIL125W | KGD1 | α-ketoglutarate dehydrogenase | • | • | -2.0 | × |
| YDR148C | KGD2 | α-ketoglutarate dehydrogenase | • | - | -2.2 | • |
| YOR142W | LSC1 | succinyl-CoA ligase | • | × | -2.5 | × |
| YGR244C | LSC2 | succinyl-CoA ligase | • | -2.8 | -3.2 | • |
| YKL148C | SDH1 | succinate dehydrogenase | × | -2.8 | -3.7 | × |
| YLL041C | SDH2 | succinate dehydrogenase | • | -2.7 | -2.4 | • |
| YKL141W | SDH3 | succinate dehydrogenase | • | -2.5 | -2.2 | • |
| YDR178W | SDH4 | succinate dehydrogenase | • | × | × | × |
| YPL262W | FUM1 | fumarate hydratase | × | -3.4 | -3.1 | × |
| YKL085W | MDH1 | malate dehydrogenase (mitochondrial) | + | -2.4 | -2.9 | • |
| Glyoxylate Cycle | | | | | | |
| YCR005C | CIT2 | citrate synthase (peroxisomal) | -6.4 | +2.1 | + | +4.7 |
| YER065C | ICL1 | isocitrate lyase | -4.1 | +2.3 | • | +9.3 |
| YNL117W | MLS1 | malate synthase | -7.1 | + | • | × |
| YOL126C | MDH2 | malate dehydrogenase | -2.2 | + | • | • |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

The level of gene expression changes in *S. cerevisiae* fermenting Icwine juice in comparison to gene expression changes in *S. cerevisiae* fermenting dilute Icwine juice are shown as either positive (+ fold upregulation) or negative (-fold downregulation) values. A plus sign (+) or minus sign (-) alone indicates significant difference in gene expression but less than a 2-fold change; • indicates no significant difference in gene expression; × indicates no expression was detected.

fructose-1,6-bisphosphate. The difference in *PFK1* expression between fermentation conditions was nearly six-fold on days three and four. Pyruvate kinase (*PYK1*) catalyzes the final and irreversible reaction of glycolysis, which involves the transfer of energy stored in phosphoenolpyruvate to ADP to form ATP. *PYK1* was upregulated on day two of the Icewine juice fermentation but was downregulated approximately three-fold on days three and four. Reduced expression of the genes involved in these three key glycolytic reactions by yeast fermenting Icewine juice also corresponded to an overall decreased rate of reducing sugar consumption throughout this fermentation, as illustrated in Figure 4.2.3.

The expression of additional genes associated with all the other steps of glycolysis and alcoholic fermentation, including those that are redox reactive, followed the same trend (Table 4.2.1). For example, Glyceraldehyde-3-phosphate dehydrogenase is encoded by three highly homologous genes (*TDH1*, *TDH2* and *TDH3*) and the gene product participates in the reversible redox reactive conversion of glyceraldehyde-3-phosphate to 1,3-bisphosphoglycerate, resulting in the formation of NADH. The expression of all three isoforms was strongly reduced up to 6.7-fold on days three and four in the Icewine juice condition relative to the dilute juice condition.

Pyruvate metabolism represents a central branch point in carbohydrate metabolism in yeast since it can either: (1) follow a reductive route, where it is decarboxylated to acetaldehyde and in turn be further reduced to ethanol by alcoholic fermentation; or (2) follow an oxidative route to form oxaloacetate and/or acetyl-CoA, both of which can enter the TCA cycle (Pronk *et al.*, 1996) (Figure 4.2.4). The fermentative route consists of two enzymatic reactions, the first of which is catalyzed by

pyruvate decarboxylase. *PDC1*, *PDC5* and *PDC6* encode for three isozymes of pyruvate decarboxylase, which catalyze the decarboxylation of pyruvate to acetaldehyde. All three *PDC* isoforms were expressed approximately two-fold higher in those yeast cells fermenting Icewine juice on day two. On days three and four, these isoforms were increasingly downregulated in the Icewine juice condition. By day five, the difference in expression of the major isoform (*PDC1*) was ten-fold between fermentations.

The second step of alcoholic fermentation is the reduction of acetaldehyde to ethanol, which regenerates oxidized cofactor, NAD^+ . This reaction is catalyzed by alcohol dehydrogenases and the cytosolic isoforms (*ADH1*, *ADH2* and *ADH5*), together with the mitochondrial isoform (*ADH3*), were for the most part downregulated throughout the first few days of the Icewine fermentation (Table 4.2.1). A minor mitochondrial isoform encoded by the gene *ADH4* showed a two-fold increase in expression in yeast fermenting Icewine juice on day two, but no expression of this gene was detected in either fermentation condition on subsequent days. The higher expression of all three *PDC* isoforms on day two of the Icewine juice fermentation, combined with the general downregulation of *ADH* expression, possibly corresponds to the spike in acetaldehyde levels measured early on in the Icewine fermentation, along with lowered ethanol levels measured at the end of the fermentation (see Section 4.5.1.1).

4.2.1.3.2. TCA and glyoxylate cycles

RNA expression signals for the genes involved in the TCA cycle were detected using the microarrays (Table 4.2.1), allowing for the expression profile of these genes to be compared between fermentation conditions. In general, the expression of genes associated with the TCA cycle was at least two-fold lower on days three and four during

the Icewine juice fermentation condition compared to the dilute juice fermentation. Two of the five genes that encode for subunits of the pyruvate dehydrogenase enzyme complex (*PDA1* and *PDB1*), which catalyzes the conversion of pyruvate to acetyl-CoA and the formation of NADH, were weakly downregulated in Icewine juice fermenting cells. The two genes encoding for both subunits of NAD⁺-dependent isocitrate dehydrogenase (*IDH1* and *IDH2*), which is responsible for catalyzing the oxidative decarboxylation of isocitrate to α -ketoglutarate, were differentially expressed between fermentation conditions. The TCA cycle intermediate α -ketoglutarate formed in this reaction then can be used for other metabolic reactions such as glutamate biosynthesis. Expression of the genes encoding for subunits of α -ketoglutarate dehydrogenase (*KGD1* and *KGD2*), succinyl-CoA ligase (*LSC1* and *LSC2*), as well as the subunits of the succinate dehydrogenase enzyme complex (*SDH1-4*), were lower at least two-fold on days three and four of the Icewine fermentation relative to the dilute juice fermentation. Malate dehydrogenase (*MDH1*) either catalyzes the final reaction of the oxidative branch of the TCA cycle, thereby producing reduced cofactor (NADH), or catalyzes the first reaction of the reductive branch, which generates oxidized cofactor (NAD⁺). The expression of this gene followed a similar pattern established by the other genes of the TCA cycle during the Icewine juice fermentation.

Despite the fact that the expression of genes encoding for enzymes of the glyoxylate cycle, such as isocitrate lyase (*ICL1*) and malate synthase (*MLS1*), have been reported in the past to be subject to glucose repression (reviewed in Krukeberg and Dickinson, 2004), expression signals were detected for these genes associated with this pathway in yeast under both fermentation conditions, even in the presence of high

concentrations glucose still remaining in the fermenting juice (see Figure 4.2.3). A 4.1- and 7.1-fold lower expression of *ICL1* and *MLS1*, respectively, was observed in Icewine juice fermenting yeast on day two compared to dilute juice fermenting yeast (Table 4.2.1). The genes encoding for peroxisomal citrate synthase (*CIT2*) and cytosolic malate dehydrogenase (*MDH2*) followed a similar trend. Aconitase (encoded by *ACO1*) is located both in the mitochondrion and in the cytosol. *ACO1* can also participate in the continuation of the glyoxylate cycle by reforming isocitrate from citrate. On day two, no significant difference in *ACO1* expression was observed between fermentations. However, increased expression levels of *CIT2* and *ICL1* were observed on days three and five of the Icewine juice fermentation.

4.2.2. Discussion - Sugar Metabolism

Glucose and fructose are the primary fermentable sugars in grape juice cofermented to generate ethanol and carbon dioxide (Fleet and Heard, 1993). Although fructose is used concurrently with glucose during table wine fermentation, *S. cerevisiae* can be described as glucophilic such that glucose is typically depleted from the juice first, giving rise to a difference between the rate of glucose and fructose consumed during fermentation (Berthels *et al.*, 2004). Under both the Icewine and dilute juice conditions, it was observed that the rate of fructose fermentation by yeast lagged behind the rate of glucose fermentation (Figure 4.2.3). However, this difference in hexose fermentation was more extensive in the Icewine juice condition. The variability between the rates of hexose utilization by *S. cerevisiae* during wine fermentation was found to be both strain dependent and is also influenced by ethanol and assimilable nitrogen levels in the must (Berthels *et al.*, 2004).

The expression profile of glycolytic and alcoholic fermentation genes from the microarray data generated in this study revealed that osmotically stressed wine yeast fermenting Icewine juice undergo a specific transcriptional change in sugar metabolism over time in response to the presence of high concentrations of fermentable sugars and this change in expression appears to slow sugar consumption rates. Sugar uptake by yeast cells has been proposed as a rate-limiting step of glycolysis (reviewed in Kruckeberg and Dickinson, 2004). Of the 20 genes encoding for different hexose transport-related proteins identified in *S. cerevisiae* (Wieczorke *et al.*, 1999), only a small subset of these were found to be differently expressed by yeast between the Icewine and diluted juice fermentation conditions (Table 4.2.1). Contrary to our results, microarray analysis has shown that *HXT5* is upregulated in laboratory yeast strains under several stress conditions including salt-induced osmotic stress (Hirayama *et al.*, 1995; Posas *et al.*, 2000; Rep *et al.*, 2000; Gasch *et al.*, 2000) and *HXT5* expression was also upregulated in wine yeast two hours after exposure to high sugar stress (Erasmus *et al.*, 2003). However, these responses are the initial responses of the yeast just hours after exposure to stress. The expression of six *HXT* genes, including *HXT5* (*HXT1*, 4, 5, 8, 10 and 12), was found to be downregulated more than two-fold in *S. cerevisiae* on days three to five of the Icewine juice fermentation. The downregulation of the *HXT* sugar transporter genes in yeast early on in Icewine fermentation may contribute to the reduced sugar consumption rate, by controlling the rate of sugar transported across the plasma membrane in the presence of over 400 g L⁻¹ of fermentable sugars in the starting juice. Also, the downregulation of these genes may also be in response to the downregulation of genes associated with glycolysis and alcoholic fermentation, since many factors affect the number of sugar

transporters occupying the plasma membrane of wine yeast cells and their affinities for hexoses. These factors include stage and rate of cell growth, nutrient availability and rate of carbon flux through the glycolytic pathway, in addition to extracellular sugar concentration (reviewed in Pretorius, 2000). The overall reduced sugar consumption rate observed in the Icewine fermentation is also in agreement with the reduction in expression of the gene encoding the rate-limiting step of glycolysis, phosphofructokinase (Table 4.2.1). Although on day two, several genes involved in the glycolytic pathway were expressed at relatively higher levels in yeast fermenting Icewine juice compared to the dilute juice, these same genes in addition to those associated with alcoholic fermentation, were significantly downregulated between fermentation conditions after day two, in agreement with the lower sugar consumption rate observed during the Icewine fermentation.

The relatively higher expression level of many of the glycolytic genes in the Icewine condition on day two of the fermentation may be related to glycerol production as opposed to ethanol production. In this study, *HXK2* and *GLK1* were significantly upregulated in wine yeast on separate days of Icewine fermentation (Table 4.2.1), but this is in opposition to the downregulation of the hexose transporters. It was previously shown by Singh and Norton (1991) that changes in yeast metabolism were induced during the adaptation of a laboratory strain of *S. cerevisiae* exposed to salt stress over an 8 h period. When exponentially growing yeast were transferred to a growth medium containing 8% (w/v) NaCl, it was found that the rates of ethanol production and glucose consumption in these stressed cells were reduced more than two-fold over those for yeast not exposed to salt. Intracellular measurements of glycolytic intermediates revealed an

accumulation of sugar phosphates associated with the upper half of the glycolytic pathway, which occurred in yeast 3 to 4 h after the transfer to salt. The decreased expression of glyceraldehyde-3-phosphate dehydrogenase (*TDH*) isoforms observed in wine yeast fermenting Icewine juice support this finding (Table 4.2.1), in that this reduced expression may separate the metabolic intermediate pools of the upper and lower halves of the glycolytic pathway and allow for the accumulation of triose phosphates, specifically DHAP, which is required for glycerol synthesis in response to osmotic stress. Both DHAP and NADH are substrates required for glycerol formation and therefore, cells need to replenish the intracellular pools of these substrates during osmoadaptation to allow glycerol synthesis to continue under hyperosmotic stress.

The increase in *PDC* expression and decrease in *ADH* expression on day two of the Icewine fermentation (Table 4.2.1) may facilitate the redirection of carbon towards acetaldehyde production for acetic acid biosynthesis and away from the more redox unfavourable pathway involving alcohol production, while osmotically stressed wine yeast cells are fermenting sugar in addition to synthesizing glycerol. Interestingly, acetaldehyde production significantly increased in the Icewine condition on day two of the fermentation (Figure 4.5.1).

Yeast cells downregulate the expression of genes involved in mitochondrial functions, including respiration and the metabolism of alternate carbon sources, such as ethanol and acetate, through carbon catabolite repression in the presence of fermentable sugars (Gancedo, 1998). It has been previously established, through genomic expression profiling using microarray analysis, that the genes associated with aerobic energy generation are induced upon a shift from glucose fermentation to respiratory growth

(Gasch *et al.*, 2000; Causton *et al.*, 2001). However, genes encoding for proteins involved in the TCA cycle, respiration and gluconeogenesis were still expressed but at very low levels during the fermentation of synthetic must containing 200 g L⁻¹ sugar (Rossignol *et al.*, 2003). In my study, the genes associated with the TCA cycle and the glyoxylate cycle were detected on the microarrays, and further, were differentially expressed between fermentation conditions (Table 4.2.1). Residual activity of TCA enzymes is maintained during fermentation to supply metabolic precursors to biosynthetic pathways (such as amino acid synthesis), but this activity results in the production of organic acids such as citrate, malate and succinate (Radler, 1993; Camarasa *et al.*, 2003). The TCA cycle has been proposed to operate in a noncyclic fashion under anaerobic conditions and can be divided into two pathways: a cytosolic reductive pathway from oxaloacetate forming NAD⁺ or FAD, and a NADH forming mitochondrial oxidative pathway beginning at α -ketoglutarate (Radler, 1993; Camarasa *et al.*, 2003). Although it is unclear what pathway operates in yeast during wine fermentation, or even if the pathways are operative, genes of the TCA cycle were expressed at significantly lower levels most prominently on day four of the Icewine juice fermentation (Table 4.2.1). So it is unlikely that either pathway (reductive or oxidative) has an impact on the cytosolic redox balance of Icewine juice cells during the first five days of fermentation. Genes associated with the glyoxylate cycle were downregulated on day two, but *CIT2* (peroxisomal citrate synthase) and *ICL1* (isocitrate lyase) were strongly upregulated as the Icewine fermentation progressed. Interestingly, the reaction catalyzed by Icl1p results in the production of both glyoxylate and succinate. So, the glyoxylate cycle may be another pathway, in addition to the oxidative and reductive pathways of the TCA cycle, which potentially contributes to

succinate production by yeast during fermentation. Under osmotic stress, the glyoxylate cycle may partially function to recycle metabolic intermediates. This pathway may also involve shuttling two carbon acetyl units in the form of acetyl-CoA between metabolites if acetyl-CoA is not being used for other anabolic reactions like amino acid synthesis (see Section 4.3), in response to a decreased availability of acetyl-CoA. The expression of the genes encoding for both acetyl-CoA synthetases (*ACS1* and *ACS2*) were found to be significantly lower in Icewine juice fermenting yeast compared to their dilute juice counterparts (refer to Figure 4.5.4) and will be further discussed in the context of acetic acid metabolism in Section 4.5.

4.3. NITROGEN METABOLISM

4.3.1. Results- Nitrogen Metabolism

4.3.1.1. Amino acid composition and nitrogen utilization

Initially, there was approximately twice the amount of assimilable nitrogen in the unfermented Icewine juice ($388.1 \pm 5.5 \text{ mg N L}^{-1}$) compared to the diluted juice ($184.7 \pm 10.7 \text{ mg N L}^{-1}$) (Figure 4.3.1A). Yeast fermenting Icewine juice also displayed a slower rate of nitrogen uptake compared to the dilute juice fermenting cells (Figure 4.3.1A).

Using the *O*-phthaldialdehyde (NOPA) assay to measure total nitrogen content in the fermenting juices, it appeared that yeast under both juice conditions only took up nitrogen during the first five days of the fermentations (Figure 4.3.1A) corresponding to entry into stationary phase. At this timepoint, dilute juice fermenting yeast cells had depleted nearly all of the pool of assimilable amino acids (Figure 4.3.1A) whereas the Icewine juice fermenting cells utilized only 20% ($78.9 \pm 2.5 \text{ mg N L}^{-1}$) of available amino acid nitrogen

present in the juice. Although the starting concentration of ammonia present in the unfermented Icewine juice was approximately two-fold higher than the concentration in the diluted Icewine juice (Figure 4.3.1B), the rate of ammonia uptake by Icewine juice fermenting cells lagged behind yeast cells fermenting the diluted juice.

The results presented here clearly show differences in both the rate and degree of total amino acid nitrogen utilization between fermentation conditions. However, information on the differences in the usage of individual amino acids by commercial wine yeast during fermentation, especially yeast under hyperosmotic stress, is very limited. The NOPA assay cannot be used to quantify individual amino acids in a wine sample. This assay is based on the derivatization of all primary amino groups to their corresponding isoindole counterparts, thereby excluding proline, in order to spectrophotometrically measure the total yeast assimilable nitrogen (YAN) levels present in a sample (Dukes and Butzke, 1998). In past studies, HPLC was used by Sanders and Ough (1985) and by Monteiro and Bisson (1991a and 1991b) to separate and quantify individual amino acids present in unfermented grape musts, using the *O*-phthaldialdehyde (OPA) derivatives of amino acids, and to determine the patterns of amino acid utilization by two commercial wine yeast strains. Therefore, in this study, HPLC was used to measure the changes in the uptake of individual amino acids, including GABA and proline by yeast during Icewine and diluted juice fermentations.

The distribution of amino acids in the unfermented Icewine juice is shown in Figure 4.3.2. Of the amino acids identified and quantified using HPLC, four amino acids, cysteine, lysine, methionine and tyrosine, were not detected in the initial Icewine and diluted juices and in the samples collected throughout the duration of the fermentations.

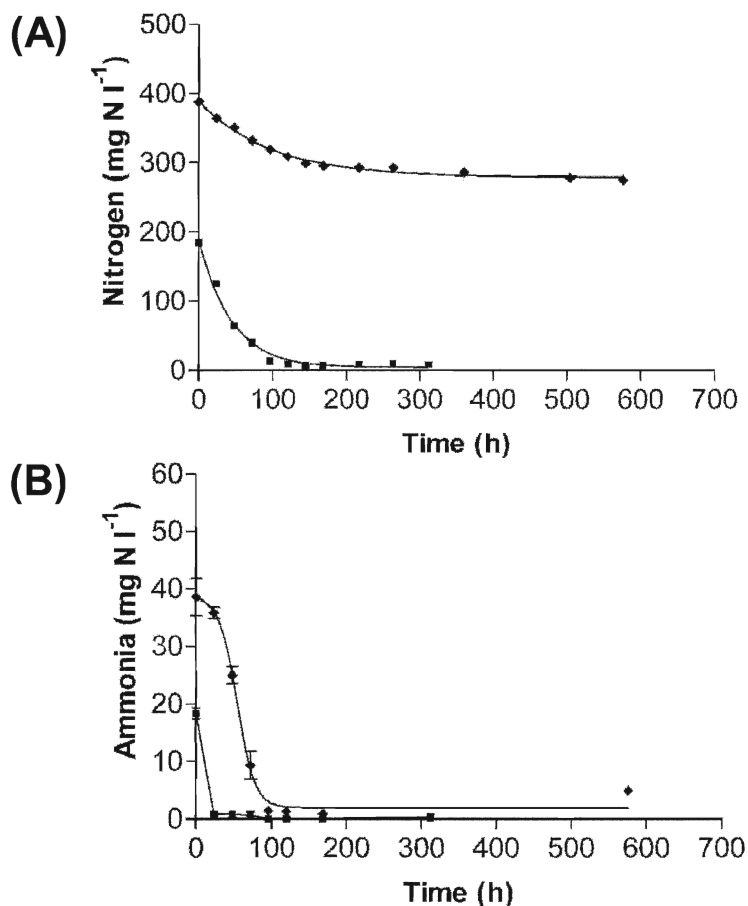


Figure 4.3.1: Yeast nitrogen utilization. Fermentations of Icewine juice (♦) and diluted Icewine juice (■) were monitored daily for total nitrogen utilization (A) and ammonia (B) throughout the course of the fermentations. Fermentations were performed in triplicate with measurements performed in duplicate. The mean values \pm standard deviation were calculated but some error bars are smaller than the symbols representing each fermentation condition.

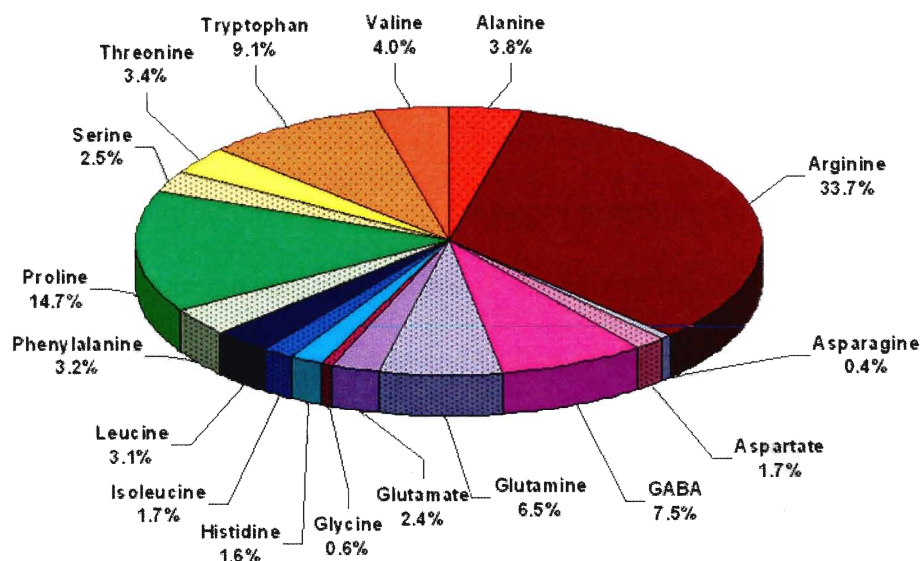


Figure 4.3.2: Distribution of amino acids in unfermented Icewine juice. The percentage of each amino acid present in the initial juice is shown.

The remaining amino acids shown in Figure 4.3.2 and listed in Tables 4.3.1 and 4.3.2 were quantified using HPLC. Although at the start of the fermentations, the diluted Icewine juice contained half the amount of each amino acid in comparison to the undiluted and unfermented Icewine juice, the percent contribution of each amino acid to the total YAN content was approximately equivalent between juices (Tables 4.3.1 and 4.3.2). Arginine and proline were the predominant amino acids present in Icewine juice. The initial arginine concentration of 1535 mg L^{-1} ($122.8 \text{ mg N L}^{-1}$) represented 33.7% of the total amino acids measured, while the contribution of the initial proline concentration of 668 mg L^{-1} to the overall amino acid content in the juice was 14.7% at the start of the fermentation. However, proline does not contribute to YAN under anaerobic conditions. Considerable concentrations of GABA, tryptophan and glutamine were also present in the starting Icewine juice, such that each nitrogen compound was found to be 7.5% and 9.1% and 6.5% of the YAN nitrogen in the juice, respectively (Table 4.3.1). Branched chain amino acids leucine, isoleucine and valine occupied a smaller portion of the total amino acids measured (less than 10% combined). Glycine made up only 0.6% of the amino acids measured. In addition to the amino acids measured, ammonia also made up a considerable portion of the YAN (nearly 10%).

Regarding the distribution of amino acid nitrogen present in the Icewine juice 168 h post-inoculation where amino acid uptake ceased, the bulk of the amino acids remaining were arginine, proline, tryptophan and GABA (Figure 4.3.3). This timepoint (168 h) was used to compare the differences in amino acid usage, as it represents the stage where yeast cells under both fermentation conditions have almost ceased nitrogen uptake and it also represents the time before cells lyse and release their intracellular

Table 4.3.1: Amino acids present in unfermented Icewine juice and finished wine.

| Amino Acid | Juice | | | Wine | | |
|------------------|--------------------|----------------------|-------------|--------------------|----------------------|-------------|
| | mg L ⁻¹ | mg N L ⁻¹ | % Total YAN | mg L ⁻¹ | mg N L ⁻¹ | % Total YAN |
| Alanine | 173.6 ± 16.4 | 27.3 ± 2.6 | 6.6 | 113.5 ± 4.0 | 17.8 ± 0.6 | 6.4 |
| Arginine | 1535.2 ± 20.1 | 122.8 ± 1.6 | 29.5 | 1273.1 ± 29.0 | 101.9 ± 10.3 | 36.6 |
| Asparagine | 20.35 ± 1.7 | 2.1 ± 0.2 | 0.5 | 13.5 ± 1.5 | 1.4 ± 0.2 | 0.5 |
| Aspartate | 76.5 ± 6.5 | 8.0 ± 0.7 | 1.9 | 48.9 ± 4.6 | 5.1 ± 0.5 | 1.8 |
| GABA | 343.1 ± 13.0 | 46.7 ± 1.8 | 11.2 | 319.6 ± 49.1 | 43.5 ± 6.7 | 15.6 |
| Glutamate | 109.6 ± 9.2 | 10.4 ± 0.9 | 2.5 | 95.7 ± 11.0 | 9.1 ± 1.0 | 3.3 |
| Glutamine | 298.1 ± 22.5 | 28.3 ± 2.1 | 6.8 | 202.9 ± 20.0 | 19.3 ± 1.9 | 6.9 |
| Glycine | 29.5 ± 1.2 | 5.5 ± 0.2 | 1.3 | 14.3 ± 1.5 | 2.7 ± 0.3 | 1.0 |
| Histidine | 72.6 ± 4.1 | 6.5 ± 0.4 | 1.6 | 45.3 ± 6.6 | 4.1 ± 0.6 | 1.5 |
| Isoleucine | 78.7 ± 7.0 | 8.4 ± 0.8 | 2.0 | 0.0 ± 0.0 | 0.0 ± 0.0 | 0.0 |
| Leucine | 139.7 ± 5.6 | 15.0 ± 0.6 | 3.6 | 23.6 ± 7.9 | 2.5 ± 0.8 | 0.9 |
| Phenylalanine | 145.6 ± 18.0 | 12.4 ± 1.5 | 3.0 | 109.6 ± 10.8 | 9.3 ± 0.9 | 3.3 |
| Proline | 668.2 ± 15.1 | - | - | 674.0 ± 57.0 | - | - |
| Serine | 114.9 ± 5.3 | 15.3 ± 0.7 | 3.7 | 54.0 ± 5.6 | 7.2 ± 0.7 | 2.6 |
| Threonine | 152.7 ± 7.4 | 18.0 ± 0.9 | 4.3 | 68.7 ± 19.7 | 8.1 ± 2.3 | 2.9 |
| Tryptophan | 416.2 ± 18.6 | 28.7 ± 1.3 | 6.9 | 383.5 ± 36.2 | 26.5 ± 2.5 | 9.5 |
| Valine | 180.1 ± 14.2 | 21.6 ± 1.7 | 5.2 | 122.3 ± 16.1 | 14.7 ± 1.9 | 5.5 |
| Total amino acid | 4554.7 ± 84.4 | 377.1 ± 9.6 | | 3557.5 ± 342.0 | 273.2 ± 27.1 | |
| Ammonia | 49.2 ± 3.2 | 38.6 ± 3.2 | 9.3 | 6.2 ± 0.5 | 4.9 ± 0.5 | 1.8 |
| Total YAN | | 415.7 | 100 | | 278.1 | 100 |

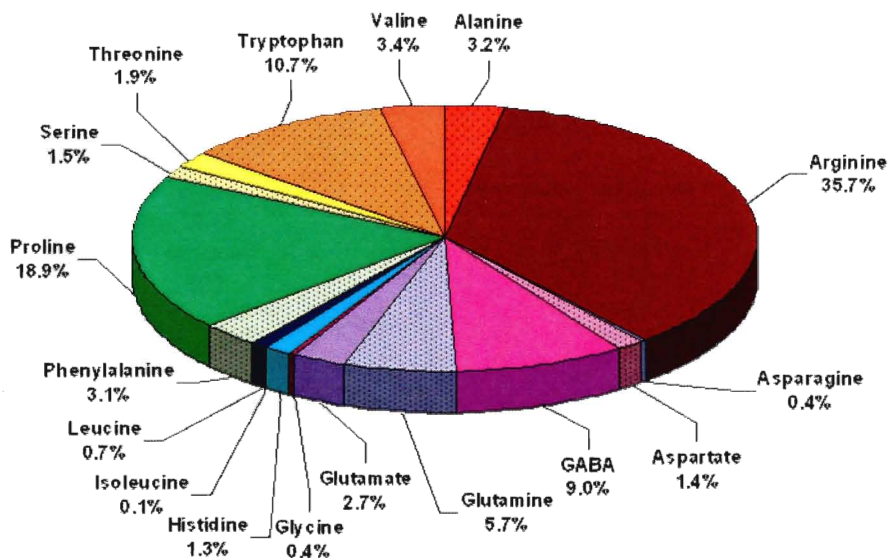
**Figure 4.3.3: Distribution of amino acids in Icewine juice 168 h post-inoculation.**

Table 4.3.2: Amino acids present in unfermented diluted Icewine juice and finished wine.

| Amino Acid | Juice | | | Wine | | |
|------------------|--------------------|----------------------|-------------|--------------------|----------------------|-------------|
| | mg L ⁻¹ | mg N L ⁻¹ | % Total YAN | mg L ⁻¹ | mg N L ⁻¹ | % Total YAN |
| Alanine | 61.2 ± 12.0 | 9.6 ± 1.9 | 5.8 | 0.6 ± 3.2 | 0.1 ± 0.5 | 2.8 |
| Arginine | 592.9 ± 34.5 | 47.3 ± 2.8 | 28.7 | 0.1 ± 2.7 | 0.0 ± 0.2 | 0.0 |
| Asparagine | 7.0 ± 0.7 | 0.7 ± 0.1 | 0.4 | 0.0 ± 0.6 | 0.0 ± 0.1 | 0.0 |
| Aspartate | 29.2 ± 2.1 | 3.1 ± 0.2 | 1.9 | 0.5 ± 0.9 | 0.1 ± 0.1 | 2.8 |
| GABA | 136.9 ± 8.2 | 18.6 ± 1.1 | 11.3 | 0.0 ± 0.0 | 0.0 ± 0.0 | 0.0 |
| Glutamate | 42.1 ± 4.2 | 4.0 ± 0.4 | 2.4 | 0.0 ± 1.8 | 0.0 ± 0.1 | 0.0 |
| Glutamine | 117.6 ± 3.8 | 11.2 ± 0.4 | 6.8 | 0.0 ± 0.0 | 0.0 ± 0.0 | 0.0 |
| Glycine | 7.4 ± 0.5 | 1.4 ± 0.1 | 0.9 | 1.0 ± 0.6 | 0.2 ± 0.1 | 5.6 |
| Histidine | 26.5 ± 3.2 | 2.4 ± 0.3 | 1.5 | 2.9 ± 1.9 | 0.3 ± 0.2 | 8.3 |
| Isoleucine | 32.0 ± 6.5 | 3.4 ± 0.7 | 2.1 | 0.0 ± 0.0 | 0.0 ± 0.1 | 0.0 |
| Leucine | 53.5 ± 2.6 | 5.7 ± 0.3 | 3.5 | 0.0 ± 0.0 | 0.0 ± 0.8 | 0.0 |
| Phenylalanine | 57.4 ± 11.2 | 4.9 ± 1.0 | 3.0 | 0.9 ± 1.9 | 0.1 ± 0.2 | 2.8 |
| Proline | 304.7 ± 16.0 | - | - | 281.9 ± 8.4 | - | - |
| Serine | 43.5 ± 2.5 | 5.8 ± 0.3 | 3.5 | 2.2 ± 0.5 | 0.3 ± 0.1 | 8.3 |
| Threonine | 58.5 ± 3.9 | 6.9 ± 0.5 | 4.2 | 4.3 ± 2.4 | 0.5 ± 0.3 | 13.9 |
| Tryptophan | 170.7 ± 14.7 | 11.8 ± 1.0 | 7.2 | 3.6 ± 0.9 | 0.3 ± 0.1 | 8.3 |
| Valine | 70.4 ± 5.5 | 8.5 ± 0.7 | 5.2 | 14.3 ± 3.3 | 1.7 ± 0.4 | 47.2 |
| Total amino acid | 1811.2 ± 98.3 | 145.3 ± 8.3 | | 310.0 ± 28.0 | 3.6 ± 2.2 | |
| Ammonia | 24.7 ± 1.7 | 19.3 ± 1.7 | 11.7 | 0.0 ± 0.0 | 0.0 ± 0.0 | 0.0 |
| Total YAN | | 164.6 | 100 | | 3.6 | 100 |

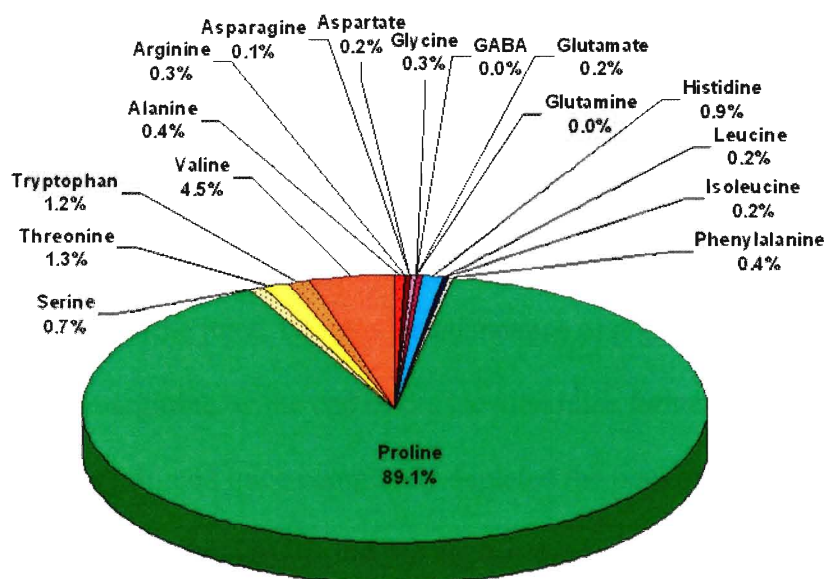


Figure 4.3.4: Distribution of amino acids in diluted Icewine juice 168 h post-inoculation.

contents into the finished wines, thereby skewing the results. Even though yeast fermenting Icewine juice failed to take up a substantial amount of each available amino acid, branched chain amino acids, leucine and isoleucine, were the only amino acids nearly or completely depleted in the juice (Table 4.3.1). In contrast, proline constituted nearly 90% of the amino acids still present in the diluted juice up to this timepoint with the residual amino acid concentrations making up the difference, individually ranging between 0-3.8% (Figure 4.3.4). In conjunction with the differences in the concentration of individual amino acids found in the finished wines compared to the starting concentrations in the initial juices, there were also obvious differences in the rate and degree of amino acid uptake over time between fermentation conditions. These results are presented in Tables 4.3.3 and 4.3.4, both of which illustrate the remaining percentage of each amino acid measured in the fermenting Icewine and diluted juices over time. Rapid uptake of amino acid nitrogen by yeast in the diluted juice condition was observed to occur 24 h into the fermentation, with near or complete depletion of the available amino acid pool by 48 to 72 h. In general, more than 50% of asparagine, aspartate, glutamine, isoleucine, leucine, serine and threonine were depleted in this juice condition after one day following inoculation. In contrast, uptake of amino acids occurred at a much slower rate by yeast fermenting Icewine juice. Despite the abundance of arginine, yeast took up only 10% of the available arginine by the end of the Icewine juice fermentation. However, yeast fermenting diluted juice completely depleted the pool of available arginine 96 h into the fermentation. GABA did not appear to be appreciably taken up by yeast fermenting Icewine juice. Conversely, by 48 h, nearly 20% of GABA was taken up

Table 4.3.3: Amino acid utilization over time by yeast fermenting Icewine juice.*

| | Time (hrs) | | | | | | |
|-------------------|------------|-------|-------|-------|-------|------|-------|
| | 0 | 24 | 48 | 72 | 96 | 120 | 168 |
| Amino Acid | | | | | | | |
| Alanine | 100.0 | 101.5 | 98.5 | 81.3 | 73.6 | 66.3 | 65.7 |
| Arginine | 100.0 | 99.9 | 97.5 | 93.5 | 90.0 | 84.3 | 82.9 |
| Asparagine | 100.0 | 97.8 | 83.5 | 74.1 | 71.7 | 59.3 | 66.4 |
| Aspartate | 100.0 | 96.2 | 88.2 | 82.5 | 76.3 | 64.8 | 64.1 |
| GABA | 100.0 | 103.0 | 98.6 | 100.1 | 102.8 | 97.3 | 93.1 |
| Glutamate | 100.0 | 103.1 | 103.5 | 100.4 | 96.9 | 89.1 | 87.4 |
| Glutamine | 100.0 | 97.2 | 88.2 | 87.9 | 77.2 | 69.6 | 68.0 |
| Glycine | 100.0 | 53.0 | 50.7 | 47.8 | 43.7 | 46.8 | 48.3 |
| Histidine | 100.0 | 96.1 | 82.9 | 79.2 | 70.5 | 62.4 | 62.4 |
| Isoleucine | 100.0 | 85.1 | 58.4 | 46.3 | 28.2 | 14.7 | 0.0 |
| Leucine | 100.0 | 84.4 | 56.7 | 44.4 | 30.6 | 19.9 | 16.9 |
| Phenylalanine | 100.0 | 93.8 | 85.3 | 88.4 | 81.2 | 69.3 | 76.0 |
| Proline | 100.0 | 104.4 | 101.5 | 102.0 | 99.3 | 99.1 | 101.0 |
| Serine | 100.0 | 95.6 | 81.1 | 73.5 | 62.5 | 49.4 | 47.1 |
| Threonine | 100.0 | 88.9 | 78.4 | 75.8 | 61.4 | 51.4 | 45.2 |
| Tryptophan | 100.0 | 106.7 | 103.8 | 93.5 | 96.0 | 88.1 | 92.0 |
| Valine | 100.0 | 97.8 | 93.0 | 85.2 | 77.8 | 72.3 | 68.0 |

* Shows the percentage of amino acids remaining in the juice relative to their starting concentrations.

Table 4.3.4: Amino acid utilization over time by yeast fermenting diluted Icewine juice.*

| | Time (hrs) | | | | | | |
|-------------------|------------|-------|-------|-------|------|------|------|
| | 0 | 24 | 48 | 72 | 96 | 120 | 168 |
| Amino Acid | | | | | | | |
| Alanine | 100.0 | 74.6 | 8.6 | 7.0 | 1.5 | 1.7 | 0.4 |
| Arginine | 100.0 | 92.9 | 37.3 | 2.2 | 0.2 | 0.4 | 0.0 |
| Asparagine | 100.0 | 22.5 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 |
| Aspartate | 100.0 | 40.0 | 0.0 | 3.2 | 1.7 | 1.7 | 1.6 |
| GABA | 100.0 | 101.4 | 82.0 | 84.1 | 28.6 | 0.0 | 0.0 |
| Glutamate | 100.0 | 76.7 | 2.8 | 0.0 | 0.0 | 0.0 | 0.0 |
| Glutamine | 100.0 | 38.2 | 0.8 | 1.2 | 0.0 | 0.0 | 0.0 |
| Glycine | 100.0 | 107.9 | 69.2 | 79.9 | 15.8 | 19.2 | 13.1 |
| Histidine | 100.0 | 77.7 | 11.1 | 3.1 | 9.7 | 12.6 | 10.6 |
| Isoleucine | 100.0 | 20.3 | 2.7 | 3.0 | 3.0 | 0.8 | 0.5 |
| Leucine | 100.0 | 22.5 | 1.1 | 0.0 | 0.0 | 1.3 | 0.0 |
| Phenylalanine | 100.0 | 89.1 | 3.3 | 3.5 | 0.3 | 1.0 | 1.4 |
| Proline | 100.0 | 110.5 | 46.9 | 51.9 | 69.4 | 72.4 | 92.6 |
| Serine | 100.0 | 22.6 | 6.0 | 6.7 | 3.5 | 5.7 | 5.0 |
| Threonine | 100.0 | 20.9 | 5.6 | 5.8 | 2.8 | 4.4 | 7.1 |
| Tryptophan | 100.0 | 114.4 | 121.2 | 126.6 | 29.0 | 5.3 | 2.1 |
| Valine | 100.0 | 79.5 | 33.6 | 36.7 | 21.9 | 24.6 | 20.2 |

* Shows the percentage of amino acids remaining in the juice relative to their starting concentrations.

by yeast fermenting diluted juice and by 120 h, all of the available GABA pool was exhausted.

With respect to the individual amino acids measured throughout the Icewine juice fermentation condition, approximately 90-100% of arginine, GABA, glutamate and tryptophan remained in the juice upon entry into stationary phase. So, 10% or less of each nitrogen compound was taken up by the yeast by the end of the fermentation. In the same fermentation condition, only 20-40% of alanine, asparagine, aspartate, glutamine, histidine and valine were taken up by yeast up to this timepoint. More than 50% of leucine, isoleucine, phenylalanine, serine, threonine and glycine were taken up by yeast by the end of the fermentation, coinciding with a lack of expression of corresponding permease genes, which will be further illustrated in subsequent sections. However, with respect to the individual amino acids measured in the diluted juice condition, there was a slight increase in glycine (less than 10%) and a more considerable increase in tryptophan (nearly 25%) over the first three days of fermentation before uptake at 96 h. There also seems to be a decrease in proline pools from days two to four even though yeast cannot use proline under fermentative conditions. Due to the large differences in nitrogen uptake between fermentation conditions, the results from the microarray experiments presented in the following section will try to correlate the changes in the expression of genes associated with nitrogen metabolism and transport to the patterns of amino acid uptake.

4.3.1.2. Gene expression profiles for nitrogen transport and metabolism

It has already been shown that Icewine juice fermenting cells consume only half of the amino acid nitrogen available to them, in comparison to dilute Icewine juice fermenting cells (Figure 4.3.1). *S. cerevisiae* is able to use different nitrogen sources for

growth but shows selectivity towards preferred nitrogen sources, like ammonia and glutamine. Repression the genes encoding for enzymes required for the utilization of poorer nitrogen sources, like ornithine, γ -aminobutyrate (GABA), allantoin and urea is invoked by a mechanism referred to as nitrogen catabolite repression (NCR) (reviewed in Magasanik and Kaiser, 2002).

The expression of yeast genes involved in nitrogen metabolism during Icewine juice fermentation was also profiled using microarray analysis (Table 4.3.5). Only four genes associated with amino acid catabolism showed significant differential expression between juice conditions. Expression of some of the genes encoding for enzymes involved in multiple amino acid biosynthesis pathways followed a similar trend. The expression of the majority of genes downregulated on day two of the Icewine juice conditions could be detected in either fermentation condition by day five.

On day two, several genes encoding for amino acid and polyamine transporters with variable substrate affinities and specificities were expressed (*AGP1*, *GNP1*, *HIP1*, *TPO2* and *TPO5*) but there was no significant difference in expression (Table 4.3.5). There was a two-fold increase in expression of the genes associated with glutamine (*GNP1*), proline (*PUT4*) and spermine (*TPO2*) transport on day three, as well as basic amino acid transport (*RHB1*) on day four of the Icewine juice fermentation. *DIP5* encodes for a dicarboxylic amino acid permease that exhibits a broad substrate specificity range, which includes glutamate (Regenberg *et al.*, 1999). *DIP5* expression was 4.7-fold lower on day two of the Icewine juice fermentation relative to the dilute juice fermentation, and corresponded to lowered expression of *GDH2* (glutamate dehydrogenase), which is involved in glutamate utilization. Expression of the GABA-

Table 4.3.5: Changes in the expression profile of genes involved in amino acid metabolism.

| ORF | Gene | Description | Fold Change* | | | |
|---|--------|--|--------------|-------|-------|-------|
| | | | Day 2 | Day 3 | Day 4 | Day 5 |
| Nitrogen Transport | | | | | | |
| YCL025C | AGP1 | broad range amino acid permease | • | • | -2.4 | • |
| YBR132C | AGP2 | broad range amino acid permease | • | • | • | × |
| YNR002C | ATO2 | ammonium transporter | -6.0 | • | • | • |
| YDR384C | ATO3 | ammonium transporter | -26.1 | × | × | × |
| YBR068C | BAP2 | branched chain amino acid permease | - | + | + | +2.0 |
| YDR046C | BAP3 | branched chain amino acid permease | -2.0 | × | × | × |
| YEL063C | CAN1 | basic amino acid permease | - | × | × | × |
| YPL265W | DIP5 | dicarboxylic amino acid permease | -4.7 | • | - | • |
| YKR039W | GAP1 | general amino acid permease | × | × | -6.0 | × |
| YDR508C | GNP1 | high affinity glutamine permease | • | +2.1 | + | × |
| YGR191W | HIP1 | basic amino acid permease | • | × | -2.5 | × |
| YNL142W | MEP2 | ammonium permease | • | - | -7.1 | × |
| YOR348C | PUT4 | proline permease | - | +2.9 | + | × |
| YCR027C | RHB1 | basic amino acid permease | + | × | +2.4 | × |
| YGR138C | TPO2 | spermine transporter | • | +2.3 | • | × |
| YOR273C | TPO4 | spermine transporter | -3.0 | × | × | × |
| YKL174C | TPO5 | putative polyamine transporter | • | -3.9 | -2.1 | × |
| YDL210W | UGA4 | GABA-specific permease | × | × | -2.1 | × |
| Amino Acid Biosynthesis and Utilization | | | | | | |
| YFL030W | AGX1 | alanine-glyoxylate transaminase | • | -3.9 | -6.9 | × |
| YOL058W | ARG1 | argininosuccinate synthase | -2.1 | × | -5.9 | × |
| YJL088W | ARG3 | ornithine carbamoyltransferase | -2.1 | +3.4 | +3.0 | × |
| YHR018C | ARG4 | argininosuccinate lyase | -2.9 | × | - | × |
| YGL202W | ARO8 | aromatic-amino-acid transaminase | + | • | -2.8 | × |
| YHR137W | ARO9 | aromatic-amino-acid transaminase | + | + | +3.2 | • |
| YDR380W | ARO10 | similar to pyruvate decarboxylase | - | -2.8 | × | × |
| YHR208W | BAT1 | branched-chain-amino-acid transaminase | - | - | • | × |
| YJR148W | BAT2 | branched-chain-amino-acid transaminase | -2.8 | -2.4 | -2.6 | × |
| YPL111W | CAR1 | arginase | +3.1 | • | • | • |
| YOR303W | CPA1 | carbamoyl-phosphate synthase | • | +4.1 | × | × |
| YAL012W | CYS3 | cystathionine gamma-lyase | +2.5 | × | +2.5 | × |
| YBR208C | DUR1,2 | urea amidolyase | • | × | × | × |
| YMR250W | GAD1 | glutamate decarboxylase | - | × | • | × |
| YOR375C | GDH1 | glutamate dehydrogenase | - | • | • | -5.3 |
| YDL215C | GDH2 | glutamate dehydrogenase | -2.5 | - | -2.2 | × |
| YAL062W | GDH3 | glutamate dehydrogenase | -3.7 | • | - | • |
| YPR035W | GLN1 | glutamine synthetase | × | - | - | -4.3 |
| YDL171C | GLT1 | glutamate synthase | - | • | - | × |
| YJR139C | HOM6 | homoserine dehydrogenase | - | × | -2.2 | × |
| YJR016C | ILV3 | dihydroxy-acid dehydratase | • | -3.0 | -3.1 | -3.0 |
| YLR355C | ILV5 | ketol-acid reductoisomerase | -2.1 | • | • | × |
| YNL104C | LEU4 | 2-isopropylmalate synthase | -5.5 | • | • | × |
| YOR108W | LEU9 | 2-isopropylmalate synthase | -5.9 | • | • | × |
| YDL182W | LYS20 | homocitrate synthase | - | + | + | +3.5 |
| YER091C | MET6 | homocysteine methyltransferase | -2.4 | • | • | × |
| YER023W | PRO3 | pyrroline-5-carboxylate reductase | • | -2.7 | -2.4 | • |
| YGR208W | SER2 | phosphoserine phosphatase | +2.4 | × | × | × |
| YER081W | SER3 | phosphoglycerate dehydrogenase | +2.7 | +2.0 | • | × |
| YDL168W | SFA1 | long-chain alcohol dehydrogenase | + | +3.1 | +3.4 | × |
| YDL080C | THI1 | alpha-ketoisocaproate decarboxylase | +2.2 | × | × | × |
| YGR019W | UGA1 | GABA transaminase | × | × | • | × |
| YBR006W | UGA2 | succinate semialdehyde dehydrogenase | × | × | • | × |
| Nitrogen Metabolism Regulation | | | | | | |
| YFL021W | GAT1 | transcription factor | +3.7 | +2.6 | +5.9 | +3.7 |
| YGL195W | GCN4 | transcription factor | • | - | -2.0 | +2.5 |
| YDL170W | UGA3 | activator of GABA catabolic genes | × | + | + | × |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

The level of gene expression changes in *S. cerevisiae* fermenting Icewine juice in comparison to gene expression changes in *S. cerevisiae* fermenting dilute Icewine juice are shown as either positive (+ fold upregulation) or negative (-fold downregulation) values. A plus sign (+) or minus sign (-) alone indicates significant difference in gene expression but less than a 2-fold change; • indicates no significant difference in gene expression; × indicates no expression was detected.

specific permease *UGA4* was approximately two-fold higher on day four of the dilute juice fermentation, corresponding to an increase of GABA uptake by these yeast cells (Table 4.3.5). However, the genes involved in both GABA synthesis (*GAD1*) and catabolism (*UGA1* and *UGA2*) were expressed at the same level between fermentation conditions also on day four. Additionally, genes encoding for ammonium ion transporters (*ATO2*, *ATO3* and *MEP2*) were similarly downregulated in the Icewine juice condition.

All pathways for nitrogen utilization converge into a common set of reactions involving glutamate and glutamine metabolism, which acts as a hub for further amino acid biosynthesis (Figure 4.3.5). Two NADP⁺-dependent glutamate dehydrogenase isoforms (*GDH1* and *GDH3*) both displayed significantly lower expression during the Icewine juice fermentation compared to the dilute juice fermentation. The enzymes encoded by these isoforms generate glutamate from ammonia and the TCA cycle intermediate α -ketoglutarate. Conversely, *GDH2* catalyzes the reverse reaction, forming NADH and NH₄⁺ in the process. *GDH2* was also significantly downregulated in yeast two-fold on days two and four of the Icewine juice fermentation. *GLN1* (glutamine synthase) and *GLT1* (glutamate synthase) are involved in the production and degradation of glutamine, respectively. *GLN1* was downregulated four-fold on day five of the Icewine juice fermentation, while *GLT1* showed a significant difference in expression between juice conditions but a less than two-fold difference change was observed. Genes encoding for some of the enzymes involved in the leucine, serine, valine and lysine biosynthetic pathways that require glutamate exhibited differential expression between juice conditions. Within the leucine biosynthetic pathway, *ILV5* (ketol-acid reductoisomerase) and *ILV3* (dihydroxy-acid dehydratase) catalyze the two step

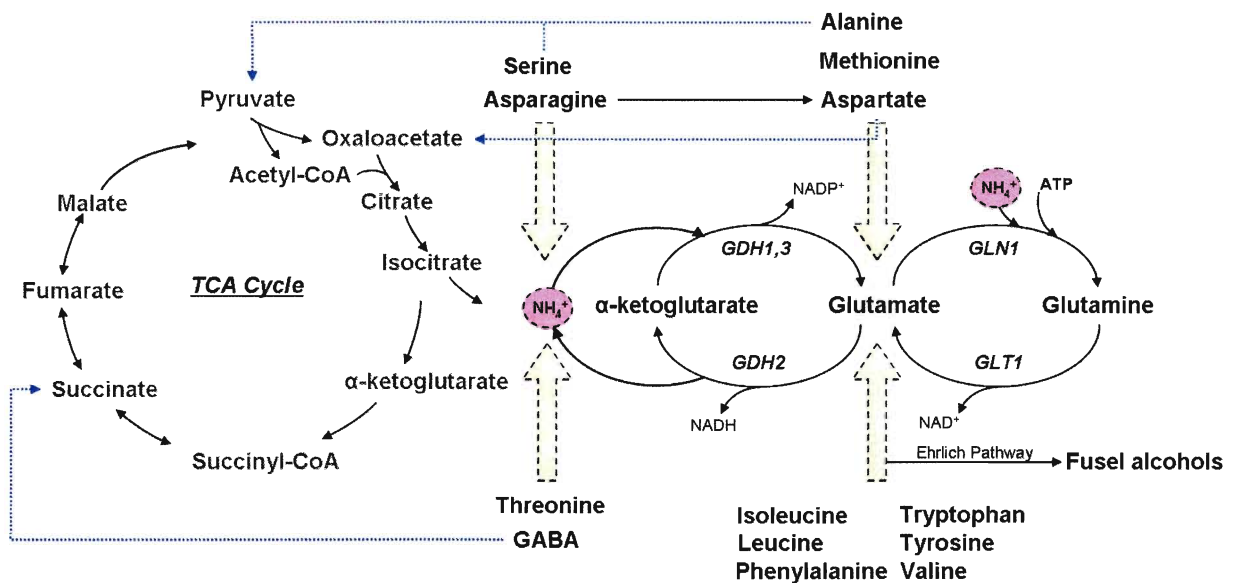


Figure 4.3.5: The central pathways of nitrogen metabolism in *S. cerevisiae*. Illustrated are the main routes associated with amino acid utilization involving deamination and transamination reactions leading to glutamate and/or ammonium ion formation (green arrows). Dashed blue lines represent pathways for the resulting carbon end products that feed into the TCA cycle. The genes associated with the enzymes involved in glutamate and glutamine metabolism are also shown (Adapted from Magasanik and Kaiser, 2002 and Godard *et al.*, 2007).

conversion of α -acetolactate to 2,3-dihydroxyisovalerate and finally to α -ketoisovalerate. Expression of both *ILV3* and *ILV5* was observed to be at least two-fold lower in Icewine juice fermenting cells. *BAT2* (branched chain amino acid transaminase) followed a similar trend (Table 4.3.5). This transaminase catalyzes the conversion of one of the intermediates of the leucine biosynthetic pathway (α -ketoisovalerate) to valine, while regenerating α -ketoglutarate from glutamate (Figure 4.3.6). *LEU4* and *LEU9* both encode for isoenzymes (2-isopropylmalate synthase) involved in the conversion of α -ketoisovalerate to α -isopropylmalate, also within the leucine biosynthetic pathway. This reaction requires the utilization of an acetyl unit from acetyl-CoA, but the expression of both *LEU4* and *LEU9* was significantly reduced five-fold on day two of the Icewine juice in comparison to the dilute juice fermentation. The serine biosynthetic pathway involves a total of three enzymatic reactions and requires glutamate, but only two of the three genes associated with this pathway were differentially expressed between juice conditions.

Alternatively, the branched chain amino acids (isoleucine, leucine and valine) can enter a common pathway and be catabolized within a series of deamination, decarboxylation and reduction reactions catalyzed by shared enzymes to form higher (fusel) alcohols, as shown in Figure 4.3.6. The enzyme encoded by *SER3* (phosphoglycerate dehydrogenase) functions to convert the glycolytic intermediate 3-phosphoglycerate to 3-phosphohydroxypyruvate, while generating NADH in the process. Expression of this gene along with the expression of *SER2* (phosphoserine phosphatase), which dephosphorylates 3-phosphoserine to form serine, were found to be higher in yeast during Icewine juice fermentation as compared to the dilute juice fermentation.

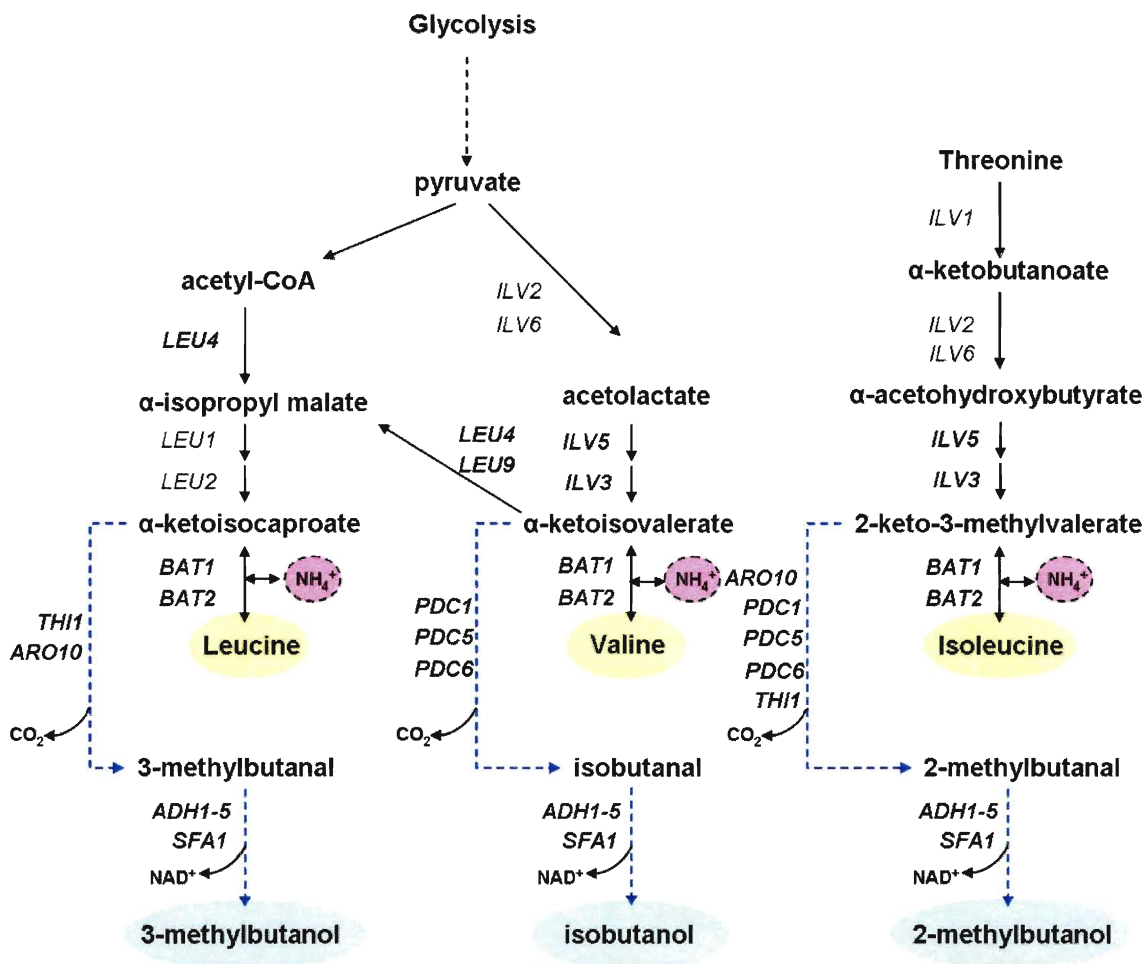


Figure 4.3.6: The pathways of branched chain amino acid metabolism in *S. cerevisiae*. Illustrated are the main routes associated with branched chain amino acid biosynthesis. Dashed blue lines represent pathways for fusel alcohol formation. Genes encoding for enzymes involved in these pathways are also shown and those in bold represent genes differentially expressed between fermentation conditions. (Adapted from Marks *et al.*, 2003).

The genes involved in arginine metabolism were also found to be differentially expressed between juice conditions (Table 4.3.5). *CARI* (arginase), which is involved in the catabolism of arginine to ornithine and urea, was expressed three-fold higher in yeast fermenting Icewine juice on day two, but no significant difference in the expression of this gene was observed between juice conditions after this timepoint. However, arginine can be synthesized from glutamine and/or ornithine directly. The expression of three genes, whose products are involved in arginine synthesis from ornithine, was approximately two-fold higher on day two of the dilute juice fermentation compared to the Icewine juice fermentation (Table 4.3.5). In this pathway, ornithine is converted to citrulline via an ornithine carbamoyl transferase (*ARG3*). Citrulline is in turn converted into L-argininosuccinate by an argininosuccinate synthase and the last reaction is catalyzed by an argininosuccinate lyase (*ARG4*), which ultimately forms arginine. Alternatively, *CPAI* encodes for a carbamoyl phosphate synthase that serves to generate carbamoyl phosphate from glutamine. This intermediate is necessary for completing the downstream reaction catalyzed by ornithine carbamoyl transferase. *CPAI* was upregulated four-fold on day three of the Icewine juice fermentation while *ARG3* was similarly upregulated on days three and four. Genes involved in the biosynthesis of aromatic amino acids (*ARO8* and *ARO9*), cysteine (*CYS3*), homoserine (*HOM6*), lysine (*LYS20*), methionine (*MET6*) and proline (*PRO3*) were also differentially expressed between fermentation conditions (Table 4.3.5).

Nitrogen metabolism is tightly regulated in *S. cerevisiae*. Both *GATI* and *GCN4* are transcriptional activators associated with the activation of a large number of NCR-sensitive genes connected with the transport and catabolism of nonpreferential amino

acids during nitrogen starvation. *GAT1* was more highly expressed throughout the Icewine juice fermentation and showed a nearly six-fold increase in expression on day four relative to the other fermentation condition (Table 4.3.5). Under Icewine fermentation conditions, *GCN4* was downregulated on days three and four but upregulated 2.5-fold on day five.

4.3.2. Discussion - Nitrogen Metabolism

4.3.2.1. Amino acid content of Icewine juice

The nitrogenous composition of Icewine juice has not been previously reported. In grape juice used for table wine production, ammonium ions and amino acids are the two main sources of yeast assimilable nitrogen compounds with ammonium ions taking up a large percentage (up to 40%) of the total YAN. Although proline and arginine are the most common nitrogenous compounds present in juice (35-60% of total amino acid content) (Beltran *et al.*, 2004; reviewed in Boulton *et al.*, 1998), proline is normally not metabolized during fermentation and must be excluded from YAN calculations. The concentrations of ammonium, proline and arginine measured in Icewine juice used in this study did fall into these projected ranges, however, it was also found that the aromatic amino acid tryptophan contributed to nearly 10% of the total amino acid content. Furthermore, like GABA, high levels of tryptophan present in the Icewine juice may be a result of subjecting the grapes from which this juice was derived to multiple freeze-thaw cycles thereby concentrating and eventually changing the juice's nitrogen profile over time.

The results presented in this study has demonstrated that despite the abundance of amino acids, osmotically stressed yeast in Icewine juice struggled to utilize both good

and poor sources of nitrogen throughout the fermentation compared to their diluted juice fermenting counterparts. In the following sections, these differences in amino acid utilization by yeast between fermentation conditions will be related to differences in the expression of genes associated with amino acid transport and metabolism.

4.3.2.2. General amino acid transport

The microarray transcription profiles of three general amino acid permeases (*AGP1*, *AGP2* and *GAP1*) revealed that these genes were expressed in yeast under Icewine and diluted juice conditions up to day five of the fermentations. In general, there was no significant difference in the expression of broad range amino acid permeases *AGP1* and *AGP2* between yeast exposed to high osmotic conditions and in the presence of non-limiting nitrogen sources compared to those under less osmotically stressful but increasingly nitrogen-limiting conditions. The expression of *AGP2* was previously observed to be downregulated by salt and sorbitol induced osmotic stress (Lee *et al.*, 2002). Also, the expression level of *AGP2* was found to be 10-fold lower than the more highly expressed general permeases (*GAP1* and *AGP1*) in laboratory yeast strains grown under nitrogen limiting conditions on minimal media (Schreve and Garrett, 2004). Despite these observations, *AGP2* was still expressed in wine yeast under both fermentative conditions in my study. *GAP1* also encodes a general amino acid permease that governs the uptake of all naturally occurring L-amino acids and other nitrogenous compounds including polyamines and GABA. But, its expression is subject to NCR thereby facilitating the transport of poorer sources in the absence of optimal nitrogen sources like glutamate and glutamine (Stanbrough and Magasanik, 1995; Regenberg *et al.*, 1999; reviewed in Boulton *et al.*, 1998; ter Schure *et al.*, 2000). In my study, it was

observed that the expression of *GAP1* was only detected on day four of the fermentations but it was expressed at much lower levels in the Icewine condition in conjunction with *AGPI*. The relatively higher expression of *GAP1* in the dilute juice condition versus the Icewine juice condition corresponds to a timepoint when nitrogen levels in the diluted juice had become extremely limiting and preferable amino acids had been completely depleted. In comparison, Icewine juice fermenting cells still had access to preferable nitrogen sources like glutamate and glutamine up to this timepoint. The lower expression of genes encoding for broad specificity amino acid transporters like *AGPI* and *GAP1* may indicate a necessity for yeast to have a tighter control over the rate of amino acid transport into the cell under Icewine fermentation conditions where high concentrations of amino acids, in addition to sugar and other solutes, contribute to the overall hyperosmotic environment. In a study by Norbeck and Blomberg (1998), uptake of methionine, leucine and histidine was found to be greatly reduced in laboratory yeast strains grown in the presence of sodium chloride compared to strains grown without salt. They proposed that reduced amino acid uptake in salt stressed yeast cells was due to a lack of the expression of high affinity amino acid transporter-encoding genes and/or by the dissipation of the proton gradient across the plasma membrane. Amino acid uptake in yeast is generally driven by proton motive force (reviewed in Boulton *et al.*, 1998) and dissipation of the proton gradient may negatively impact yeast cells' ability to import amino acids. This latter point will be further discussed in Section 4.4 in the context of the glycerol/H⁺ symporter *STL1* which was found to be highly expressed during Icewine fermentation. Exposure to high osmotic stress and expression of low activity and low

affinity bulk amino acid transporters like *AGP2*, may also contribute to the overall reduction in amino acids taken up by yeast fermenting Icewine juice.

4.3.2.3. Specific amino acid uptake and metabolism

The branched chain amino acids (leucine, isoleucine and valine) combined contributed to less than 10% of the total amino acid content in the initial Icewine juice. Interestingly, of all the amino acids present, only leucine and isoleucine were completely depleted from this juice at the end of the fermentation. Uptake of leucine and isoleucine by yeast fermenting Icewine juice corresponded to a relative increase in *BAP2* expression up to day five. *BAP2* along with its isoform *BAP3* are closely related to *AGP1* and are capable of transporting branched chain and aromatic amino acids (Regenberg *et al.*, 1999). It is unclear why yeast fermenting Icewine juice would favour the uptake of leucine and isoleucine in the presence preferential nitrogen sources (glutamate and glutamine) whose concentrations remained relatively high in the juice until the end of the fermentation. However, the microarray data showed a small subset of genes associated with leucine and isoleucine biosynthesis that were either downregulated in yeast fermenting Icewine juice relative to those fermenting dilute juice or were expressed, but showed no significant difference between fermentation conditions. If not being actively synthesized by yeast, the influx of leucine and isoleucine transported into the cells may be diverted to other pathways such as protein synthesis or fusel alcohol production via the Ehrlich pathway.

During all fermentations, small amounts of higher (fusel) alcohols can be formed from the transamination products of branched chain and aromatic amino acids. However, fusel alcohol synthesis usually occurs during the later stages of fermentation following

the period of rapid amino acid uptake (reviewed in Boulton *et al.*, 1998). During fusel alcohol synthesis, amino acids are first transaminated to their corresponding α -ketoacids and in turn, α -ketoglutarate accepts an ammonia group to form glutamate (Figure 4.3.6). In the second step, the α -ketoacid is decarboxylated to produce an aldehyde. The final reaction of this redox reactive pathway involves the reduction of the aldehyde by an alcohol dehydrogenase and formation of a fusel alcohol. This pathway allows nitrogen in the form of ammonia to be retrieved (Dickinson, 2004). Dickinson *et al.* (2003) presented evidence that the final step of fusel alcohol formation via the conversion of an aldehyde to a fusel alcohol could be performed by any of the five *ADH* genes and *SFA1* (formaldehyde dehydrogenase). Although not measured in the fermenting juices, fusel alcohols may have been formed early on in the Icewine fermentation as a consequence of the upregulation of the pyruvate decarboxylases (*PDC1,5* and *6*) (Table 4.2.1) along with the decarboxylase encoded by *THI1* and the dehydrogenase encoded by *SFA1* (Table 4.3.5). If Icewine fermenting yeast are able to produce fusel alcohols, these cells would have likely been limited at the beginning of this pathway due to the relative downregulation of the transaminase encoding genes *BAT1* and *BAT2*. Furthermore, since this pathway is redox reactive, the production of NAD^+ in a cell which may already be in cytosolic redox imbalance, would potentially place additional stress on yeast cells fermenting Icewine juice.

Fusel alcohol production is one of many pathways involving the recycling of an ammonia group from another amino acid to form glutamate. Both glutamate and glutamine are important amino group donors in the biosynthesis of amino acids and other nitrogen containing compounds (reviewed in Boulton *et al.*, 1998; Magasanik and Kaiser,

2002). It has been shown in this study that Icewine juice fermenting cells fail to take up a large proportion of the available glutamate and glutamine present in the juice. A small number of glutamate and glutamine-specific transporters have been identified in yeast. *DIP5* encodes for a glutamate transporter which is also capable of transporting glutamine, aspartate and asparagine (Regenberg *et al.*, 1999). From the microarray data, *DIP5* was not found to be expressed at the same degree in Icewine juice fermenting cells as compared to dilute juice fermenting cells. *GNP1* encodes for a transporter capable of transporting glutamine (Regenberg *et al.*, 1999) but was found to be more highly expressed in yeast during Icewine fermentation. Glutamate and glutamine transport is also mediated by *GAP1* (Regenberg *et al.*, 1999). Generally, the slower rate of uptake of glutamate and glutamine by Icewine fermenting yeast cells corresponds to an overall downregulation of genes encoding for carriers responsible for their transport. These amino acids may also be competing with each other for available transporters potentially driving down their ability to be taken up by the cell even further. A decrease in glutamate and glutamine uptake also corresponds to the downregulation of glutamate metabolism genes. The formation of glutamate from α -ketoglutarate involves the enzymatic activity of glutamate dehydrogenases encoded by *GDH1* and *GDH3* (Figure 4.3.5). The expression of *GDH1* and *GDH3* was reported to be dependent on carbon source by DeLuna *et al.*, (2001). Gdh3p activity was found to be glucose repressed in yeast but activity increased on non-fermentable carbon sources like ethanol whereas Gpdh1p activity contributed to glutamate formation during exponential growth and in the presence of high glucose concentrations. However, expression of both *GDH* isoforms during Icewine and diluted juice fermentation seem to be independent of the availability

of glucose. There are two routes for the catabolic production of glutamate from glutamine via enzymes encoded by *GDH2* or *GLT1/GLN1* (Figure 4.3.5). However, in the presence of glutamate and glutamine, both *GDH1* and *GDH2* are subject to NCR (reviewed in ter Schure *et al.*, 2000). *GDH3*, and to a lesser extent *GDH2*, were upregulated in wine yeast cells briefly exposed to high sugar stress (Erasmus *et al.*, 2003). This was not the case in our study since yeast were placed under prolonged exposure to hyperosmotic stress during Icewine fermentation. The depletion of both glutamate and glutamine in the diluted juice condition, together with the probable lifting of NCR by day two of the fermentation, could account for the increase expression of these genes compared to yeast fermenting Icewine juice.

Proline and arginine were the predominant amino acids measured in Icewine juice. Yeast cannot use proline as a source of nitrogen in anaerobic conditions (reviewed in Boulton *et al.*, 1998). Under aerobic conditions, yeast can degrade proline to glutamate since the first enzymatic step in the proline catabolic pathway occurs in the mitochondrion and is catalyzed by a proline oxidase (encoded by *PUT1*). The second and last step of the proline degradation pathway is catalyzed by a dehydrogenase (*PUT2*) (reviewed in Boulton *et al.*, 1998). Proline utilization is also subject to NCR (reviewed in Boulton *et al.*, 1998; Magasanik and Kaiser, 2002). As expected, proline was not utilized or metabolized by yeast fermenting either Icewine or diluted Icewine juices and the microarray data supports these observations since none of the genes involved in proline catabolism were expressed under either fermentation condition. In addition to proline, *PUT4* can facilitate the import of alanine and glycine (Regenberg *et al.*, 1999) and GABA (Andre *et al.*, 1993). Although *PUT4* expression is regulated transcriptionally by

NCR thereby preventing proline utilization if better nitrogen compounds are present (ter Schure *et al.*, 2000), this gene was found to be expressed in Icewine juice fermenting cells up to three-fold higher than that found in the dilute condition. *PUT4* upregulation could be correlated to an increased uptake of glycine by cells in Icewine juice compared to those in diluted juice. But, increased glycine uptake may be associated with amino acid transporters other than *PUT4*. Elevated *PUT4* expression may also be attributed to the hyperosmotic stress imposed on yeast cells fermenting Icewine juice as this gene was reported to be upregulated by both brief high salt- and sugar-induced hyperosmotic stress (Posas *et al.*, 2000; Rep *et al.*, 2000; Erasmus *et al.*, 2003). Proline uptake by diluted juice fermenting yeast may be a result of cells being relieved of NCR as the fermentation progressed but they were unable to use it as a nitrogen source under anaerobic conditions.

Arginine transport is mediated by *GAP1* in addition to the arginine-specific permease *CAN1* (Regenberg *et al.*, 1999). Downregulation of both transporters by Icewine juice fermenting cells may have led to an overall reduction in arginine uptake. Despite these observations the expression of *CAR1*, which encodes an NCR-sensitive arginase responsible for degrading arginine to ornithine and urea, was significantly higher on day two in the Icewine juice condition. In contrast, three genes associated with arginine biosynthesis (*ARG1*, 3 and 4) were concurrently downregulated. The higher expression of *CAR1* in yeast during Icewine fermentation may be in response to hyperosmotic stress since this gene was strongly upregulated more than 25-fold in sorbitol-stressed laboratory yeast strain (Rep *et al.*, 2000). The activity of urea amidolyase (encoded by *DUR1,2*) can further metabolize urea to ammonia and carbon dioxide. The microarray results show that *DUR1,2* was expressed only on day two but

there was no significant difference in expression between fermentation conditions. Since urea was not measured in the fermenting juices in this study it would be impossible to conclude if increased expression of *CAR1* and lack of expression of *DUR1,2* by Icewine fermenting cells lead to increased urea levels in the wine. Ornithine can be further degraded to form polyamine precursors of spermine, spermidine and putrescine (reviewed in Boulton *et al.*, 1998). Although the genes involved in spermine, spermidine and putrescine biosynthesis (*SPE1-4*) were not expressed in yeast fermenting either Icewine or diluted juice the polyamine exporters (*TPO2*, 4 and 5) were downregulated in Icewine fermenting cells. Polyamines are required for yeast growth (reviewed in Boulton *et al.*, 1998) however it is unclear whether wine yeast utilize polyamines during fermentation, especially those under Icewine fermentation conditions where yeast growth is limited.

GABA also made up a considerable percentage of the available nitrogen measured in Icewine juice as expected, since GABA is among the major nitrogen sources present in grape juice (Monteiro and Bisson, 1991a). GABA transport in yeast is mediated by the GABA-specific permease *UGA4* (Andre *et al.*, 1993) in addition to *GAP1* and *PUT4*. No appreciable GABA appeared to be taken up by Icewine fermenting yeast. Typically, yeast do not use all available GABA due to the effects of NCR (Monteiro and Bisson, 1991b). However, yeast fermenting diluted juice did uptake GABA but only after approximately 120 mg N L⁻¹ of amino acid nitrogen had already been taken up from the juice. This possibly indicated that cells were being relieved of NCR. An increase in *GAP1* and *UGA4* expression in conjunction with lifting of NCR may be correlated to a substantial increase in GABA uptake after day three by yeast in

the diluted juice condition where nearly all the amino acids had already been depleted from the fermenting juice. *UGA1* encodes a 4-aminobutyrate aminotransferase that deaminates GABA to succinate semialdehyde which in turn, is converted to succinate by the NADP⁺-dependent succinate semialdehyde dehydrogenase (SSADH) encoded by *UGA2* (Ramos *et al.*, 1985). In the presence of good nitrogen sources, *UGA1* is also subject to NCR (reviewed in ter Schure *et al.*, 2000). As seen in Table 4.5.1, there was no significant difference in the concentration of succinate found in the finished wines. Since the genes involved in the GABA degradation pathway were generally not expressed in yeast under both juice conditions, it is unlikely that this pathway contributes to succinate production during the first five days of the fermentations. Day four of the fermentations seems to be a key timepoint for GABA transport and metabolism. Although the genes involved in its synthesis and degradation were expressed on day four, no significant difference was found between fermentation conditions. In the presence of GABA, expression of *UGA1* and *UGA2* is induced which is mediated by the transcriptional activator Uga3p (Vissers *et al.*, 1989). Interestingly, *UGA3* was upregulated in the Icewine juice condition. Expression of *GAD1* and *UGA2* are also upregulated under conditions of oxidative stress (Coleman *et al.*, 2001). In 2003, Erasmus *et al.* also found that exposure to brief sugar stress upregulated the expression of GABA metabolism genes in wine yeast. Even though Icewine juice fermenting cells took up approximately 7% of the available GABA present in the juice, expression of genes associated with GABA metabolism may be attributed to the general stress response by yeast. Previous studies have shown that GABA may act as a compatible osmolyte in plants subjected to salt stress (reviewed in Shelp *et al.*, 1999). Based on the gene profiling data and amino acid

usage results presented in our current study, it is unlikely that *S. cerevisiae* uses GABA for osmoregulation during Icewine juice fermentation.

4.4. GLYCEROL METABOLISM

4.4.1. Results- Glycerol Metabolism

4.4.1.1. Glycerol Production

The concentration of extracellular glycerol measured during both the Icewine and diluted juice fermentations was found to increase over time, yet there was little difference in the rate of glycerol production by yeast as a function of time between fermentation conditions over the first 120 h (Figure 4.4.1A). Following this timepoint, glycerol production subsequently continued to increase over time in the Icewine juice fermentation resulting in a final glycerol concentration of $11.95 \pm 0.16 \text{ g L}^{-1}$, while the rate of glycerol production plateaued in the dilute juice fermentation leading to a final glycerol concentration of $6.30 \pm 0.19 \text{ g L}^{-1}$ (Table 4.4.1). This represents a significant 1.9-fold increase in glycerol production by yeast fermenting Icewine juice compared to the dilute juice after subtracting the initial glycerol concentration in the starting juices. Since the rates of sugar consumption differed greatly between fermentation conditions (Figure 4.2.3) and glycerol is a byproduct of sugar metabolism, the amount of glycerol produced was plotted against sugar consumed to compare the rate of glycerol production as a function of sugar consumed (Figure 4.4.1B). Although Icewine juice fermenting yeast consumed less sugar over time, more glycerol was released by these cells throughout the fermentation after an equal amount of sugar was consumed in comparison to the dilute juice fermenting yeast.

Table 4.4.1: Glycerol produced by *S. cerevisiae* following the consumption of 187 g L⁻¹ of sugar.

| Fermentation | Initial reducing sugar in juice* (g L ⁻¹) | Final reducing sugar in wine* (g L ⁻¹) | Total sugar consumed (g L ⁻¹) | Glycerol produced* (g L ⁻¹) |
|----------------------|---|--|---|---|
| Icewine juice | 447.7 ± 8.5 | 259.9 ± 2.8 | 187.8 ± 2.8 | 11.95 ± 0.16 |
| Dilute Icewine juice | 226.7 ± 4.1 | 39.9 ± 3.1 | 186.8 ± 3.1 | 6.30 ± 0.19 |

* Values within the same column are significantly different as determined by a paired t-test ($P \leq 0.05$)

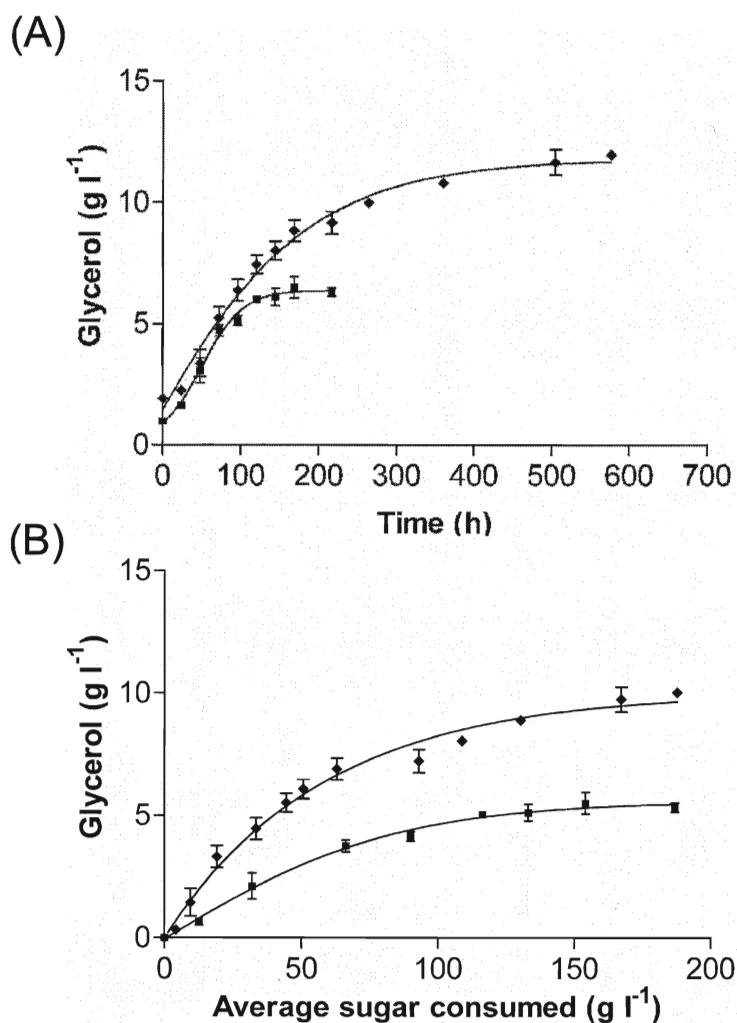


Figure 4.4.1: Yeast glycerol production. The concentration of extracellular glycerol (A) was measured in duplicate throughout the course of the fermentations of Icewine juice (♦) and diluted Icewine juice (■) and was plotted as a function of sugar consumed (B). The values represent the average ± standard deviation of the mean of triplicate fermentations.

4.4.1.2. Gene expression profiles (Microarrays)- Glycerol metabolism and transport

The genes involved in glycerol synthesis were generally more highly expressed in yeast fermenting Icewine juice. Both *GPD1* and *GPD2* encode for NAD⁺-dependent glycerol-3-phosphate dehydrogenase isoenzymes that catalyze the first redox reactive and rate-limiting step in glycerol synthesis (Larsson *et al.*, 1993; Eriksson *et al.*, 1995; Remize *et al.*, 2001). However, only *GPD1* showed significant differential expression between fermentation conditions (Table 4.4.2). *GPD1* was upregulated four-fold on day three, but the fold difference in the expression of this gene between the Icewine and dilute juice conditions peaked on day four (6.6-fold). Although *GPD2* was found to be expressed during days two and three of the fermentations, there was little to no significant difference in its expression between juice conditions. By day five, the expression of both *GPD* isoforms could not be detected using the cDNA microarrays for all three replicate fermentations. Both glycerol-3-phosphatase isoforms (*GPP1* and *GPP2*) were observed to have similar expression patterns throughout the Icewine juice fermentation (Table 4.4.2). *GPP1* and *GPP2* were strongly expressed in the Icewine condition relative to the dilute juice condition. The fold change in the expression of *GPP1* and *GPP2* between fermentation conditions peaked on day four. After which, the level of upregulation for these genes began to decline by day five in comparison to the dilute juice fermentation. Also seen in Table 4.2.2, two genes encoding for enzymes responsible for catalyzing the conversion of glycerol back into the glycolytic intermediate DHAP (*GUT1* and *GUT2*) were found to be expressed at the same level between fermentation conditions on day two. However, the expression of both genes was significantly lower by 2.4-fold on day three of the Icewine juice fermentation compared

to the dilute juice fermentation. Increased expression of glycerol synthesis genes along with decreased expression of glycerol catabolism genes during the early stages of Icewine juice fermentation corresponds to increased levels of glycerol produced by these yeast (Figures 4.4.1A and B).

Accumulation of glycerol under hyperosmotic stress is also dependent on the yeast cell's ability to retain it within the cytoplasm. *FPS1* encodes for a glyceroporin channel that passively facilitates the export of glycerol (Van Aelst *et al.*, 1991; Luyten *et al.*, 1995; Sutherland *et al.*, 1997; Oliveira *et al.*, 2003). *FPS1* was expressed at the same level between fermentation conditions on day two (Table 4.4.2). After day two, the expression of *FPS1* could not be detected in yeast fermenting either juice. In contrast, *STL1* which encodes for a putative glycerol/H⁺ symporter (Ferreira *et al.*, 2005), was one of the most strongly upregulated genes identified in yeast fermenting Icewine juice (Table 4.2.2). This was intriguing since the expression of *STL1* was previously found to be transiently induced under salt stress (Rep *et al.*, 2000) and subject to glucose repression (Ferreira *et al.*, 2005). Throughout the Icewine juice fermentation, the fold change in *STL1* expression increased and ultimately peaked on day five (25.6-fold) as compared to the dilute juice fermentation.

Table 4.4.2: Expression profile of the genes involved in glycerol metabolism.

| ORF | Gene | Description | Fold Change [*] | | | |
|-----------------|------|------------------------------------|--------------------------|-------|-------|-------|
| | | | Day 2 | Day 3 | Day 4 | Day 5 |
| Glycerol | | | | | | |
| YDL022W | GPD1 | glycerol-3-phosphate dehydrogenase | + | +4.3 | +6.6 | × |
| YOL059W | GPD2 | glycerol-3-phosphate dehydrogenase | + | • | × | × |
| YIL053W | GPP1 | glycerol-3-phosphatase | +3.1 | +8.7 | +12.3 | +3.5 |
| YER062C | GPP2 | glycerol-3-phosphatase | +3.0 | +9.3 | +14.0 | +4.7 |
| YHL032C | GUT1 | glycerol kinase | • | -2.4 | -5.4 | × |
| YIL155C | GUT2 | glycerol-3-phosphate dehydrogenase | • | -2.4 | × | × |
| YLL043W | FPS1 | glycerol channel | • | × | × | × |
| YDR536W | STL1 | glycerol proton symporter | +11.5 | +11.9 | +18.9 | +25.6 |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

The level of gene expression changes in *S. cerevisiae* fermenting Icewine juice in comparison to gene expression changes in *S. cerevisiae* fermenting dilute Icewine juice are shown as either positive (+ fold upregulation) or negative (-fold downregulation) values. A plus sign (+) or minus sign (-) alone indicates significant difference in gene expression but less than a 2-fold change; • indicates no significant difference in gene expression; × indicates no expression was detected.

4.4.1.3. Gene expression profiles (Northern)- Glycerol

From the microarray data, the expression profile of the gene *STL1* (glycerol/H⁺ symporter) was shown to be highly upregulated in yeast fermenting Icewine juice. Since this gene displayed a greater than ten-fold change in expression between fermentation conditions, it was decided that the expression of *STL1* in yeast under Icewine and diluted juice fermentation conditions required validation using Northern analysis. However, it was necessary to validate the specificity of the DNA probe designed exclusively for *STL1*.

Salt induced hyperosmotic stress was used for *STL1* probe validation since *STL1* had been previously shown to be strongly induced in laboratory yeast strains exposed to saline stress (Rep *et al.*, 2000). Figure 4.4.2 shows that under salt stress the *STL1* probe was specific to detect the expression of *STL1* in the commercial wine yeast strain of *S. cerevisiae* K1-V1116 in addition to the laboratory strain BY4742. A weak signal was detected in the *STL1* deletion mutant (Δ *STL1*) under inductive conditions, however, this signal was 17- and 11-fold lower than the signal generated from either the wine yeast or the laboratory yeast strains, respectively. This signal may be attributed to background noise from the blot as there was no visible band observed on the Northern.

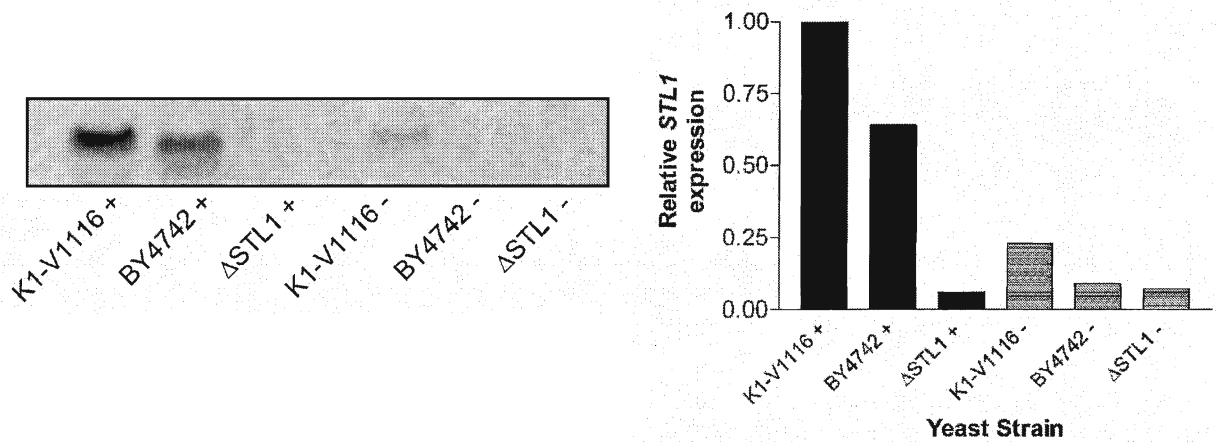


Figure 4.4.2: Probe specificity for *STL1* validation Northern blots. Yeast cultures were induced (+) or not induced (-) with salt prior to RNA isolation. Expression profiles of the blot following hybridization of 32 P-labeled DNA probes specific for *STL1* to RNA isolated from the commercial wine yeast strain (K1-V1116), parent yeast strain (BY4742) and knockout yeast strain (Δ STL1).

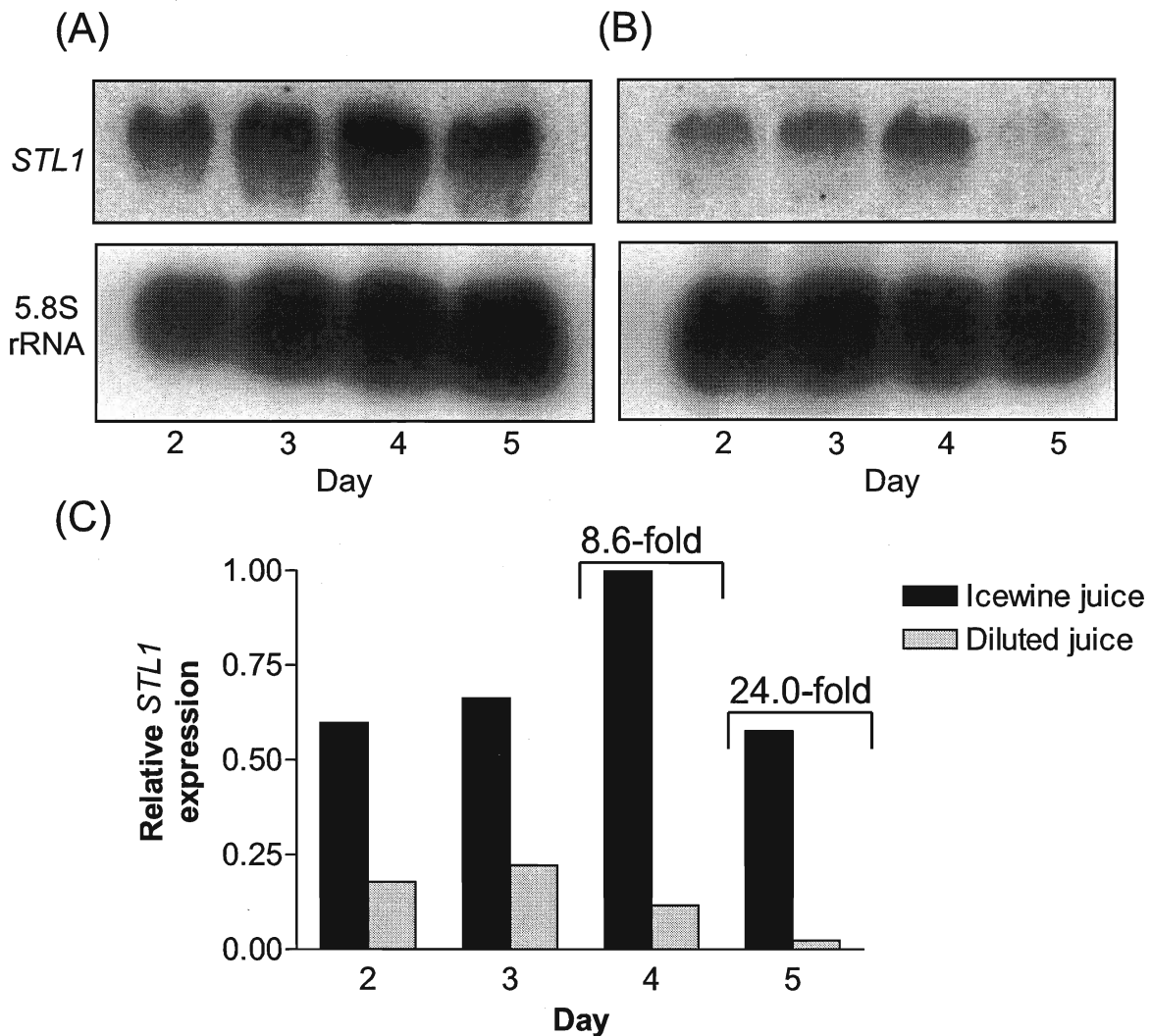


Figure 4.4.3: Expression of *STL1* during fermentation. The expression profiles of *STL1* from yeast fermenting Icewine juice (A) and diluted Icewine juice (B) using Northern analysis. 5.8S ribosomal RNA was used as a loading control. (C) Comparison of the relative levels of *STL1* expression between fermentation conditions.

The relative gene expression profile of *STL1* during Icewine and dilute wine fermentation is depicted in Figure 4.4.3. From the Northern blot, *STL1* was expressed throughout both fermentation conditions but to a much greater extent in yeast fermenting Icewine juice. The signal intensity for *STL1* peaked on day four of the Icewine juice fermentation in comparison to the signal intensity of the dilute juice fermentation, representing a 8.6-fold difference in expression. However, the relative fold difference in *STL1* expression peaked on day five between fermentation conditions giving rise to a 24-fold increase in *STL1* expression in Icewine juice fermenting cells similar to that found with the microarrays.

4.4.2. Discussion - Glycerol metabolism and transport

The combination of Gpd1p-Gpp2p activity is proposed to be mainly involved in glycerol production during osmotic stress, while the Gpd2p-Gpp1p combination is connected with balancing the NAD^+/NADH cofactor system under anaerobic conditions (Albertyn *et al.*, 1994b; Ansell *et al.*, 1997; Pålman *et al.*, 2001). Wine yeast face both anaerobic and osmotic stress during Icewine fermentation, so either or both of these factors may control the expression of genes associated with glycerol synthesis.

At the start of fermentation, glycerol is produced in response to the excess NADH generated during yeast cell biomass formation and its synthesis acts to restore redox balance by reoxidizing reduced cofactor (van Dijken and Scheffers, 1986). However, the fermentation of Icewine juice results in a reduction in biomass formation but an increase in glycerol production. The microarray results obtained in the present study showed that the expression of *GPD2* did not significantly differ between Icewine and dilute juice fermentation conditions and therefore *GPD2* does not appear to be contributing to the

increased glycerol levels found in the Icewine fermentation. Since *GPD2* expression would not be required for glycerol production to restore the cytosolic pool of NAD^+ since biomass formation is diminished, the elevated production of glycerol in wine yeast during Icewine fermentation corresponded to an upregulation of *GPD1* expression. These microarray results are in agreement with studies by Pigeau and Inglis (2005 and 2007), in which the increase in *GPD1* expression as measured by Northern analysis, but not *GPD2* expression, over time in wine yeast fermenting Icewine juice compared to dilute juice corresponded to elevated glycerol levels produced throughout the fermentation. From our study, it appears that glycerol synthesis is also dependent on both glycerol-3-phosphatase isoforms during Icewine juice fermentation as *GPP1* and *GPP2* were observed to be strongly expressed. The expression of *GUT1* (glycerol kinase) and *GUT2* (glycerol-3-phosphate dehydrogenase) is subject to glucose repression and does not appear to be induced by osmotic stress (reviewed in Hohmann, 2002). However, both glycerol catabolizing genes were more highly expressed on day three of the dilute juice fermentation in which over 160 g L^{-1} of fermentable sugars remained in the juice up to this timepoint. The osmotic stress imposed on yeast by high concentrations of sugars is dynamic since the extracellular osmotic pressure exerted on cells decreases as they take up and eventually ferment the sugar. An increase in *GUT1* and *GUT2* may be a reflection of the decreasing osmotic stress in the dilute juice condition and a lower demand for high intracellular glycerol levels by yeast fermenting this juice in comparison to the Icewine juice fermenting cells.

To ensure that the glycerol produced in *S. cerevisiae* upon exposure to hyperosmotic stress is accumulated to offset the osmotic pressure exerted on the plasma

membrane, yeast must regulate glycerol export through the plasma membrane. *FPS1* encodes for a glyceroporin channel that functions as a glycerol exporter. Once yeast cells return to normal growth conditions following osmotic shock, Fps1p facilitates passive transport of glycerol across the membrane down a concentration gradient (Van Aelst *et al.*, 1991; Luyten *et al.*, 1995; Sutherland *et al.*, 1997; Oliveira *et al.*, 2003). However, this channel rapidly closes when yeast are exposed to hyperosmotic stress (Támas *et al.*, 1999), thereby controlling the amount of glycerol retained by the cell. No expression of this gene could be detected in yeast fermenting either Icewine or diluted juice after day two. Although measurable, there was no significant difference in the expression of this gene between fermentation conditions. Despite the lack of *FPS1* expression, extracellular glycerol levels measured in the fermenting juices increased steadily over the days subject to microarray analysis. It is possible that *FPS1* expression was below the detection level of microarray analysis, or a number of glyceroporin channels had already occupied the yeast plasma membrane early on during the fermentations such that *FPS1* expression was no longer required after day two. It is also likely that the glycerol released by yeast cells into the fermenting juices occurred by simple diffusion across the plasma membrane. However, rate of glycerol diffusion across the membrane is generally low and membrane permeability to glycerol is reduced even further in osmotically stressed yeast cells (Tamás *et al.*, 1999; Sutherland *et al.*, 1997).

Active glycerol uptake via a glycerol/H⁺ symporter (Stl1p) may be occurring in *S. cerevisiae* during Icewine juice fermentation. *STL1* was one of the most strongly expressed genes identified in yeast fermenting Icewine juice. Expression of this gene was previously found to be transiently induced in laboratory yeast strains under salt stress

(Rep *et al.*, 2000) and is reported to be subject to glucose repression (Ferreira *et al.*, 2005). However, in agreement with our study, *STL1* was one of the most strongly upregulated genes in wine yeast following a two hour exposure to high sugar stress (Erasmus *et al.*, 2003). A comparison of the expression profiles of this gene throughout the Icewine juice and dilute juice fermentations using both microarray and Northern analysis showed a considerable differential increase in *STL1* expression in yeast fermenting Icewine juice. This ultimately resulted in a peak 25-fold increase in *STL1* expression on day five of the fermentation relative to the level in the dilute juice condition. The roles of Fps1p and Stl1p in glycerol transport in the presence of high concentrations of fermentable sugars during Icewine fermentation are unclear. Osmotolerant yeast species like *Zygosaccharomyces rouxii* are capable of actively importing glycerol against a concentration gradient into the cell in the presence of salt-induced hyperosmotic stress (van Zyl *et al.*, 1990). Active glycerol uptake is a mechanism yeast can use to increase intracellular glycerol concentrations under hyperosmotic stress and it has been suggested that a high affinity glycerol/H⁺ symporter uptake system also exists in *S. cerevisiae* (Lages and Lucas, 1997). The use of Stl1p by yeast for glycerol uptake during Icewine juice fermentation may contribute to limited cell growth by dissipating the proton gradient across the plasma membrane in an energy dependent manner. Many yeast amino acid transport systems are proton symports (reviewed in Boulton *et al.*, 1998; Horák, 1986) therefore any disturbance in the proton gradient across the cell membrane may negatively influence amino acid uptake. It has been shown in Section 4.3 that the rate and degree of amino acid uptake by yeast fermenting Icewine juice is far lower than that of dilute juice fermenting cells. However

if yeast cells are using Stl1p to actively take up glycerol in the Icewine juice condition, then the increased influx of protons into the cytoplasm may ultimately hinder active amino acid uptake. The ability to excrete excess protons is an important regulatory factor for amino acid uptake in yeast in addition to preventing cytoplasm acidification and cell death (reviewed in Boulton *et al.*, 1998).

4.5. ACETIC ACID METABOLISM

4.5.1. Results - Acetic Acid Metabolism

4.5.1.1. Acetic Acid Production

Since acetaldehyde is the precursor to acetic acid, extracellular acetaldehyde and acetic acid levels were both measured throughout the fermentations. The rate of acetaldehyde production increased rapidly in the Icewine juice fermentation and reached a maximum concentration of $0.13 \pm 0.01 \text{ g L}^{-1}$ in 96 h (Figure 4.5.1A). This corresponds to a 5.5-fold difference in acetaldehyde concentration between juice conditions at this timepoint. Acetaldehyde levels then quickly declined eventually leading to a significant 1.5-fold increase in acetaldehyde concentration at the end of the Icewine juice fermentation in comparison to the end of the fermentation of the diluted juice. Similar trends were observed when the amount of acetaldehyde released into the juice was plotted against sugar consumed (Figure 4.5.1B).

The rate of extracellular acetic acid released into the fermenting juices greatly differed between fermentation conditions (Figure 4.5.1C). Acetic acid was produced and released at a faster rate and to a greater extent during the first five days of the Icewine juice fermentation in comparison to the dilute juice condition. The rate slowed down and

Table 4.5.1: Metabolites produced by *S. cerevisiae* following the consumption of 187 g L⁻¹ of sugar.

| Fermentation | Acetaldehyde* (g L ⁻¹) | Acetic Acid* (g L ⁻¹) | Succinate (g L ⁻¹) | Ethanol* % (v/v) |
|----------------------|---------------------------------------|--------------------------------------|-----------------------------------|---------------------|
| Icewine juice | 0.05 ± 0.00 | 1.40 ± 0.03 | 0.68 ± 0.05 | 11.5 ± 0.8 |
| Dilute Icewine juice | 0.03 ± 0.00 | 0.14 ± 0.01 | 0.64 ± 0.01 | 14.4 ± 0.2 |

* Values within the same column are significantly different as determined by a paired t-test ($P \leq 0.05$)

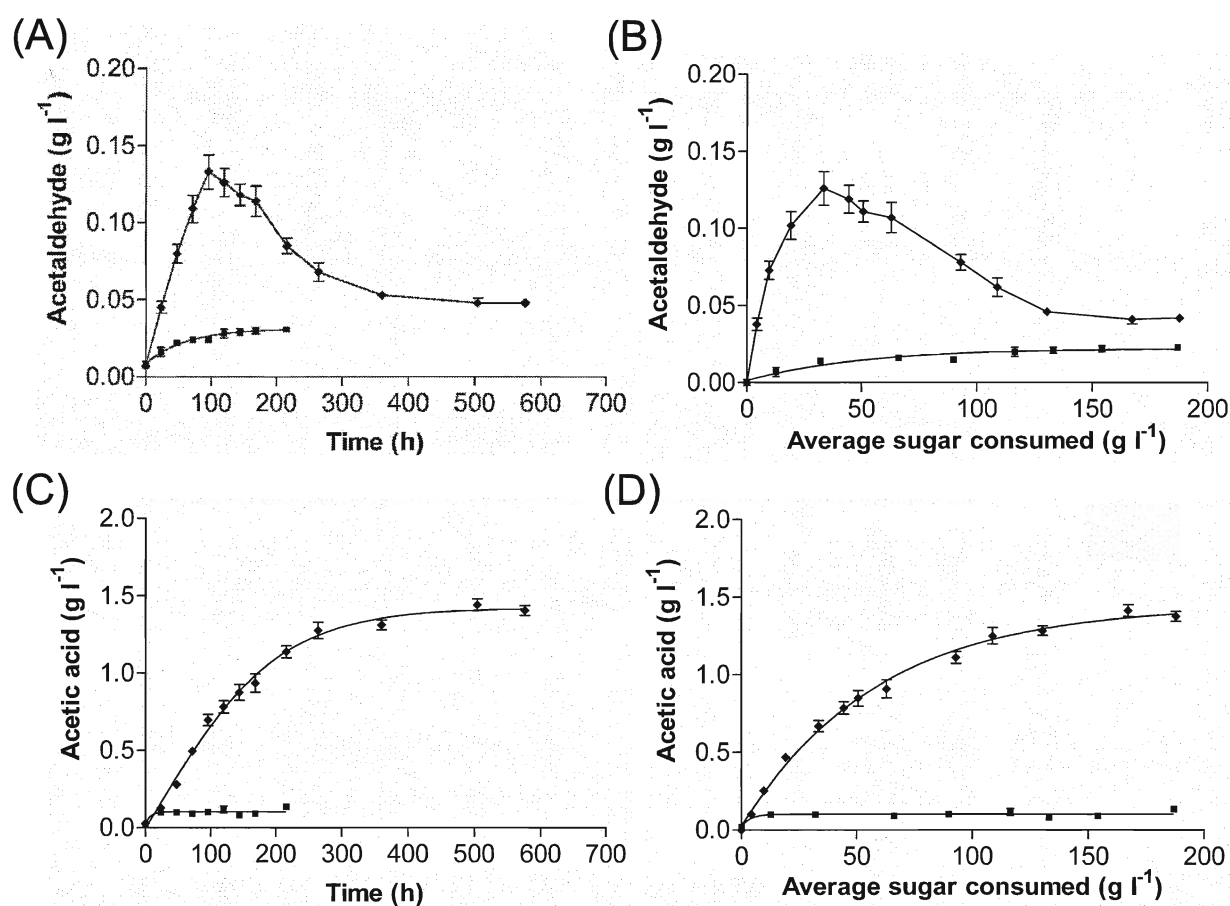


Figure 4.5.1: Yeast acetaldehyde and acetic acid production. Acetaldehyde (A) and acetic acid (C) were measured in duplicate throughout the course of the fermentations of Icewine juice (♦) and diluted Icewine juice (■). Acetaldehyde (B) and acetic acid (D) were plotted as a function of sugar consumed. The values represent the average ± standard deviation of the mean of triplicate fermentations.

eventually plateaued after 300 h of fermentation resulting in a final extracellular acetic acid concentration of $1.40 \pm 0.03 \text{ g L}^{-1}$ (Table 4.5.1). In contrast, acetic acid was produced at a slower rate up to 48 h into the dilute juice fermentation with the extracellular acetic acid concentration remaining constant at $0.14 \pm 0.01 \text{ g L}^{-1}$ until the end of the fermentation. Wine yeast fermenting Icewine juice produced 10-fold more acetic acid than yeast fermenting the diluted juice after approximately 187 g L^{-1} of sugar was consumed (Figure 4.5.1.D).

In contrast to the measured levels of glycerol, acetaldehyde and acetic acid produced by Icewine juice fermenting yeast, these cells produced significantly less ethanol than the dilute juice cells by the end of the fermentations when 187 g L^{-1} of sugar was consumed (Table 4.5.1). A decrease in ethanol production by osmotically stressed yeast cells fermenting Icewine juice may indicate a change in the metabolic flux of hexose sugars through glycolysis and alcoholic fermentation. Carbon derived from sugar metabolism is possibly being diverted away from ethanol synthesis and towards the production of other metabolites for necessary survival and adaptation to hyperosmotic stress.

4.5.1.2. Gene expression profiles for acetic acid and fatty acid metabolism

Of the five genes encoding for aldehyde dehydrogenases identified in *S. cerevisiae*, only *ALD3* encoding for a cytosolic NAD^{+} -dependent aldehyde dehydrogenase was differentially expressed for three of the four fermentation days examined (Table 4.5.2). *ALD3* was strongly expressed on day two of the Icewine juice fermentation relative to the dilute condition. The differential expression of this gene continued to increase as the fermentation progressed whereupon the relative fold change

Table 4.5.2: Expression profile of the genes involved in acetic acid metabolism.

| ORF | Gene | Description | Fold Change [*] | | | |
|--------------------|------|--|--------------------------|-------|-------|-------|
| | | | Day 2 | Day 3 | Day 4 | Day 5 |
| Acetic acid | | | | | | |
| YAL054C | ACS1 | acetyl-CoA synthetase | -19.0 | • | • | × |
| YLR153C | ACS2 | acetyl-CoA synthetase | -4.8 | -3.6 | -11.2 | × |
| YBL015W | ACH1 | acetyl-CoA hydrolase | -5.2 | -2.3 | -2.9 | +2.4 |
| YCR010C | ADY2 | acetate transporter | -13.9 | +3.2 | +5.6 | +14.1 |
| YMR170C | ALD2 | aldehyde dehydrogenase | • | • | +2.2 | × |
| YMR169C | ALD3 | aldehyde dehydrogenase | +8.4 | +9.7 | +14.6 | × |
| YOR374W | ALD4 | aldehyde dehydrogenase (mitochondrial) | -6.2 | • | • | • |
| YER073W | ALD5 | aldehyde dehydrogenase (mitochondrial) | -4.4 | • | • | × |
| YPL061W | ALD6 | aldehyde dehydrogenase | -2.4 | + | • | • |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

The level of gene expression changes in *S. cerevisiae* fermenting Icewine juice in comparison to gene expression changes in *S. cerevisiae* fermenting dilute Icewine juice are shown as either positive (+ fold upregulation) or negative (-fold downregulation) values. A plus sign (+) or minus sign (-) alone indicates significant difference in gene expression but less than a 2-fold change; • indicates no significant difference in gene expression; × indicates no expression was detected.

in *ALD3* expression peaked on day four (14.6-fold) of the Icewine juice fermentation. In contrast both mitochondrial *ALD* isoforms (*ALD4* and *ALD5*) along with the cytosolic NADP⁺-dependent isoform (*ALD6*) were expressed up to six-fold lower in yeast fermenting Icewine juice relative to the dilute condition (Table 4.5.2). The expression level of these three genes was not significantly different between fermentation conditions after day two. *ALD2* encodes for an additional cytosolic NAD⁺-dependent aldehyde dehydrogenase and was found to be transiently upregulated in yeast on day four of the Icewine juice fermentation. *ADY2* encodes for a protein that has been proposed to function as an acetate permease (Paiva *et al.*, 2004). *ADY2* expression was found to be far lower (13.9-fold) on day two of the Icewine fermentation relative to the dilute condition. As the fermentation progressed, however, *ADY2* expression increased over that found in the dilute condition where the fold change in expression of *ADY2* eventually peaked on day five (14.1-fold).

The genes encoding for two acetyl-CoA synthetases (*ACS1* and *ACS2*), which are involved in the generation of acetyl-CoA from acetic acid were two of the most strongly downregulated genes in yeast fermenting Icewine juice in comparison to the expression level in the dilute juice condition (Table 4.5.2). The mitochondrial isoform (*ACS1*) was downregulated 19-fold on day two while the cytosolic isoform (*ACS2*) was downregulated throughout the Icewine juice fermentation with the relative fold downregulation for *ACS2* peaking on day four (11-fold). Expression of both *ACS* isoforms could not be detected after day four in either fermentation condition. Acetic acid and coenzyme A (CoA) can also be regenerated through the hydrolysis of acetyl-CoA by an acetyl-CoA hydrolase (encoded by *ACH1*). The expression profile of *ACH1* was

similar to that of *ACS2*, however, *ACH1* was found to be expressed 2.4-fold higher on day five of the Icewine juice fermentation. Increased *ALD3* expression in conjunction with decreased expression of both acetyl-CoA synthetase isoforms may have contributed to the increased acetaldehyde and acetic acid levels produced by Icewine juice fermenting yeast compared to those fermenting diluted juice (Figure 4.5.1).

A considerable number of genes in *S. cerevisiae* associated with lipid and sterol metabolism, which involves the input or production of acetyl-CoA, displayed noticeably similar expression patterns in yeast under the Icewine juice condition. The expression of the vast majority of genes involved in fatty acid metabolism, phospholipid and ergosterol synthesis were relatively higher at least two-fold on day two of the dilute juice fermentation as shown in Table 4.5.3. However, as yeast continued to ferment both Icewine and diluted juices, the expression of many of these genes could not be detected in either juice condition (Table 4.5.3). In addition to fatty acids, phospholipids and sterols are important membrane components and genes involved in phospholipid and sterol metabolism were also found to be differentially expressed between fermentation conditions.

Four genes encoding for enzymes associated with phosphatidic acid metabolism showed a two-fold difference in expression (*LPP1*, *SLC1*, *AYR1* and *OPI3*). Downregulation of the ergosterol synthesis (*ERG*) genes also corresponds to the downregulation of the *ACS* (acetyl-CoA synthetase) genes. The products of the *ACS* genes can function to supply acetyl-CoA units required for ergosterol synthesis. Ergosterol is the primary sterol in yeast and one of its primary functions is to maintain

Table 4.5.3: Expression profile of the genes involved in lipid metabolism.

| ORF | Gene | Description | Fold Change* | | | |
|-------------------------|-------|--|--------------|-------|-------|-------|
| | | | Day 2 | Day 3 | Day 4 | Day 5 |
| Fatty Acid Metabolism | | | | | | |
| YML042W | CAT2 | carnitine O-acetyltransferase | -4.4 | • | × | × |
| YDR256C | CTA1 | catalase | • | × | +2.1 | × |
| YLR284C | ECI1 | dodecenoyl-CoA delta-isomerase | -4.2 | × | × | × |
| YBR026C | ETR1 | enoyl-[acyl-carrier protein] reductase | -3.1 | × | -2.5 | × |
| YIL009W | FAA3 | long-chain-fatty-acid-CoA ligase | -2.2 | × | × | × |
| YMR246W | FAA4 | long-chain-fatty-acid-CoA ligase | -2.1 | × | × | × |
| YKL182W | FAS1 | S-malonyltransferase | -4.1 | × | × | × |
| YKR009C | FOX2 | 3-hydroxyacyl-CoA dehydrogenase | -2.3 | × | × | × |
| YNL009W | IDP3 | isocitrate dehydrogenase | -2.0 | - | -2.8 | × |
| YIL160C | POT1 | acetyl-CoA C-acyltransferase | -2.1 | × | • | × |
| YGL205W | POX1 | acyl-CoA oxidase | -2.5 | × | × | × |
| YPL147W | PXA1 | fatty acid transport | -2.0 | × | × | × |
| YNL202W | SPS19 | 2,4-dienoyl-CoA reductase | -3.5 | × | × | × |
| YAR035W | YAT1 | carnitine O-acetyltransferase | -7.2 | × | × | × |
| YER024W | YAT2 | carnitine O-acetyltransferase | -4.6 | +2.7 | +2.3 | +3.1 |
| Phospholipid Metabolism | | | | | | |
| YIL124W | AYR1 | acylglycerone-phosphate reductase | -2.2 | × | - | × |
| YMR162C | DNF3 | phospholipid-translocating ATPase | -4.1 | × | × | × |
| YDR503C | LPP1 | phosphatidate phosphatase | +2.4 | × | × | × |
| YJR073C | OPI3 | methylene-fatty-acyl-phospholipid synthase | -2.1 | -2.1 | - | × |
| YNL231C | PDR16 | phosphatidylinositol transporter | +2.1 | × | × | × |
| YDL052C | SLC1 | 1-acylglycerol-3-phosphate O-acyltransferase | +2.2 | × | × | × |
| Sterol Metabolism | | | | | | |
| YGR175C | ERG1 | squalene monooxygenase | -2.1 | × | -2.0 | × |
| YML008C | ERG6 | sterol 24-C-methyltransferase | -2.5 | × | • | × |
| YPL028W | ERG10 | acetyl-CoA C-acetyltransferase | -2.1 | × | -4.3 | × |
| YHR007C | ERG11 | sterol 14-demethylase | -2.1 | -2.2 | -2.2 | × |
| YML075C | HMG1 | hydroxymethylglutaryl-CoA reductase | -3.7 | × | × | × |
| YHR001W | OSH7 | oxysterol binding | -2.0 | × | • | × |
| YDR213W | UPC2 | sterol biosynthesis transcription factor | +2.6 | × | × | × |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

The level of gene expression changes in *S. cerevisiae* fermenting Icewine juice in comparison to gene expression changes in *S. cerevisiae* fermenting dilute Icewine juice are shown as either positive (+ fold upregulation) or negative (-fold downregulation) values. A plus sign (+) or minus sign (-) alone indicates significant difference in gene expression but less than a 2-fold change; • indicates no significant difference in gene expression; × indicates no expression was detected.

membrane fluidity (Schweizer, 2004). In general, the ergosterol (*ERG*) synthesis genes found to be differentially expressed between conditions were downregulated approximately two-fold on days two and four (Table 4.5.3). *ERG10* is involved in the first reaction of the ergosterol biosynthetic pathway. *ERG10* encodes for a thiolase that catalyzes the condensation of two acetyl-CoA molecules to form acetoacetyl-CoA. Hydroxymethylglutaryl-CoA (HMG-CoA) reductase (encoded by *HMG1*) which catalyzes the reduction of HMG-CoA to mevalonate showed a four-fold lower differential expression in Icewine juice only on day two of the fermentation. Ergosterol synthesis requires oxygen and of the three other *ERG* genes (*ERG1*, *ERG6* and *ERG11*) that were less highly expressed in yeast under the Icewine juice condition, *ERG1* encodes for squalene monooxygenase which catalyzes the first oxidation reaction of the ergosterol biosynthetic pathway.

The main pathway of fatty acid degradation in *S. cerevisiae* is through β -oxidation, a process which occurs exclusively within the peroxisomes and oxygen is required for this process (reviewed in Hiltunen *et al.*, 2003) (Figure 4.5.2). The end result of the β -oxidation pathway is the formation of acetyl-CoA units which in turn can be used for biosynthetic reactions. Since the peroxisomal membrane is essentially impermeable to small molecules like cofactors and acetyl-CoA (van Roermund *et al.*, 1995), transport of larger molecules like fatty acids into peroxisomes is dependent on their conversion into acyl-CoA esters via fatty acid-CoA ligases (*FAA3* and *FAA4*). Import of these activated esters into the peroxisomal matrix is mediated by a membrane bound transporter protein encoded by the gene *PXA1*. Together, these three genes involved in fatty acid activation and transport were relatively downregulated

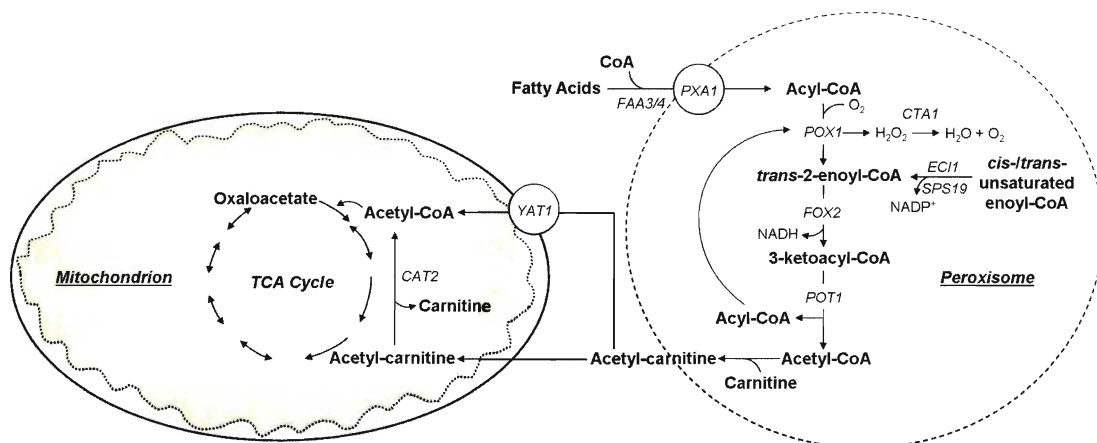


Figure 4.5.2: The central pathway of fatty acid β -oxidation in *S. cerevisiae*. Illustrated are the major metabolites generated within the main route of peroxisomal β -oxidation of fatty acids in yeast. The genes associated with the enzymes involved in fatty acid degradation are also shown. (Adapted from Hiltunen *et al.*, 2003).

approximately two-fold on day two of the Icewine fermentation (Table 4.5.3).

A large group of genes associated with β -oxidation pathway were more highly expressed in yeast on day two of the dilute juice fermentation compared to the Icewine juice fermentation (Table 4.5.3). This group included genes like *POX1*, which encodes a peroxisomal acyl-CoA oxidase that catalyzes the first (and rate-limiting) reaction of the β -oxidation process by converting the acyl-CoA ester to *trans*-2-enoyl-CoA (Figure 4.5.2). The H_2O_2 generated by this oxygen consuming reaction is removed through its decomposition to water and oxygen by a peroxisomal catalase A (*CTA1*). There was no significant difference in the expression of *CTA1* between juice conditions on day two. *ECII* (an isomerase) and *SPS19* (a $NADP^+$ -dependent reductase) both can participate in the β -oxidation of unsaturated enoyl-CoAs. Activity of the peroxisomal isocitrate dehydrogenase (*IDP3*) can regenerate $NADPH$ thereby maintaining the peroxisomal $NADP^+/NADPH$ cofactor pool. The last two enzymatic steps of the β -oxidation pathway involves *FOX2* and *POT1* which generates acetyl-CoA and a shortened acyl-CoA ester which then reenters the β -oxidation cycle.

Export of acetyl-CoA from the peroxisome is completed by its conjugation to carnitine via carnitine acetyl transferases encoded by three genes (*CAT2*, *YAT1* and *YAT2*) (Swiegers *et al.*, 2001). All three genes were more highly expressed in the dilute juice condition relative to the Icewine juice condition on day two of the fermentations. However, *YAT2* was more highly expressed in Icewine juice fermenting yeast after day two. Acetyl-CoA transported to the mitochondrion through this mechanism can enter the TCA cycle or be used towards other biosynthetic pathways.

4.5.1.3. Gene expression profiles (Northern)- Acetic acid

Along with *STL1*, the gene expression profiles of *ACSI* and *ACS2* (mitochondrial and cytosolic acetyl-CoA synthetases) generated from the microarray data were found to be highly expressed in yeast fermenting diluted Icewine juice and had also showed a greater than ten-fold change in expression between fermentation conditions. The expression profile of these two genes in addition to the NAD^+ -dependent aldehyde dehydrogenase *ALD3* was further validated using Northern analysis. The specificity of the DNA probes designed for *ACSI* and *ACS2* required validation prior to monitoring the expression of these selected genes throughout the first five days of Icewine and diluted Icewine juice fermentations. The DNA probe for *ALD3* was found to be specific for this gene and had been previously validated by Pigeau and Inglis (2005).

ACSI and *ACS2* share 66% sequence homology but induction of each gene is triggered by different metabolic conditions in yeast. Respiratory growth and glucose limitation was used to induce *ACSI* expression (Van den Berg *et al.*, 1996) while anaerobic growth conditions were used to induce *ACS2* expression (Van den Berg and Steensma, 1995). Under aerobic conditions, the *ACSI* probe was found to be specific for its target gene as the probe bound to the RNA from the ΔACS2 strain but not to the RNA from the ΔACSI strain (Figure 4.5.3A). Under anaerobic conditions, the *ACS2* probe generated a strong signal from the wine yeast strain, the parent strain and the ΔACSI strain. A weak signal for the ΔACS2 strain was detected. This weak signal may have been due to high background signal by failure to remove non-specific probe binding during the membrane washing procedure (Figure 4.5.3B). Thus, the DNA probes designed for *ACSI* and *ACS2* appear to be specific for their targets.

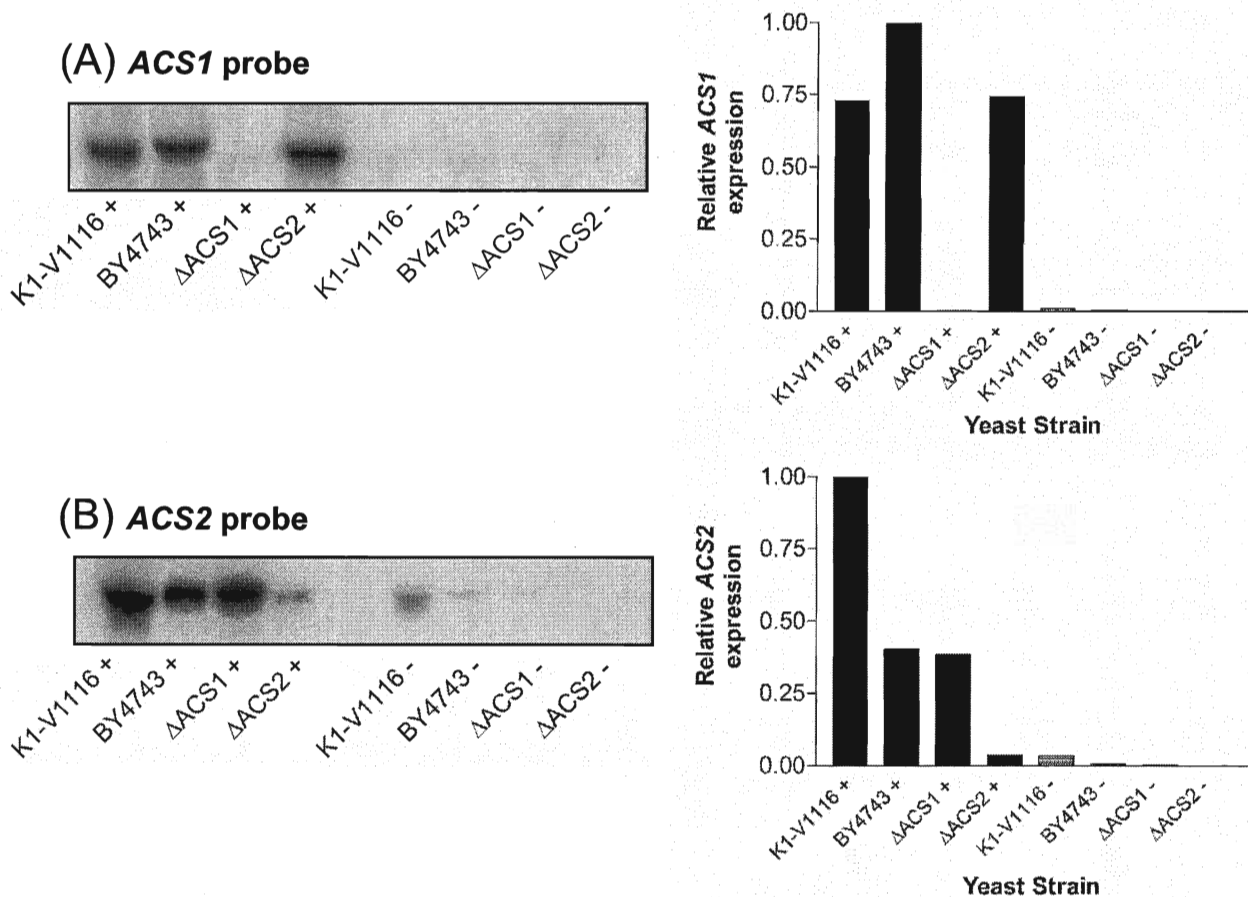


Figure 4.5.3: *ACS1* and *ACS2* probe specificity for validation Northern blots. Yeast cultures were induced to express *ACS1* under aerobic conditions (+) or not induced (-) under anaerobic conditions (A) or were induced to express *ACS2* under anaerobic conditions (+) or not induced (-) under aerobic conditions (B). Expression profiles of the blots following hybridization of ^{32}P -labeled DNA probes specific for *ACS1* and *ACS2* to RNA isolated from the commercial wine yeast strain (K1-V1116), parent yeast strains (BY4742 and BY4743) and knockout yeast strains ($\Delta ACS1$ and $\Delta ACS2$) are shown. Relative gene expression is also shown in the bar graphs.

The relative gene expression profiles for *ACSI/ACS2* and *ALD3* are depicted in Figures 4.5.4 and 4.5.5, respectively. The Northern expression profile shows that the relative signal intensity of *ACSI* was 29-fold higher in yeast fermenting the dilute juice on day two of the fermentations. As both fermentations progressed, *ACSI* was weakly expressed at approximately the same level between fermentation conditions. The relative level of *ACS2* expression increased in the dilute juice condition over time until day four and at this timepoint, a 13.2-fold difference in *ACS2* expression between fermentation conditions was observed. By day five, there was no difference in *ACS2* expression. *ALD3* was also expressed throughout both fermentations and the relative difference in peak fold expression occurred on day four of the Icewine juice fermentation giving rise to a 16-fold increase in *ALD3* expression.

The fold changes in expression for this small subset of genes determined from Northern analysis was similar to the results obtained from the microarray analysis. However, the magnitude in fold changes differed slightly between gene expression techniques but the trends were the same. Also, expression of *ACSI*, *ACS2* and *ALD3* was weak but measurable on day five of the fermentations using Northern analysis but no expression could be detected in either juice condition on this day using microarray analysis.

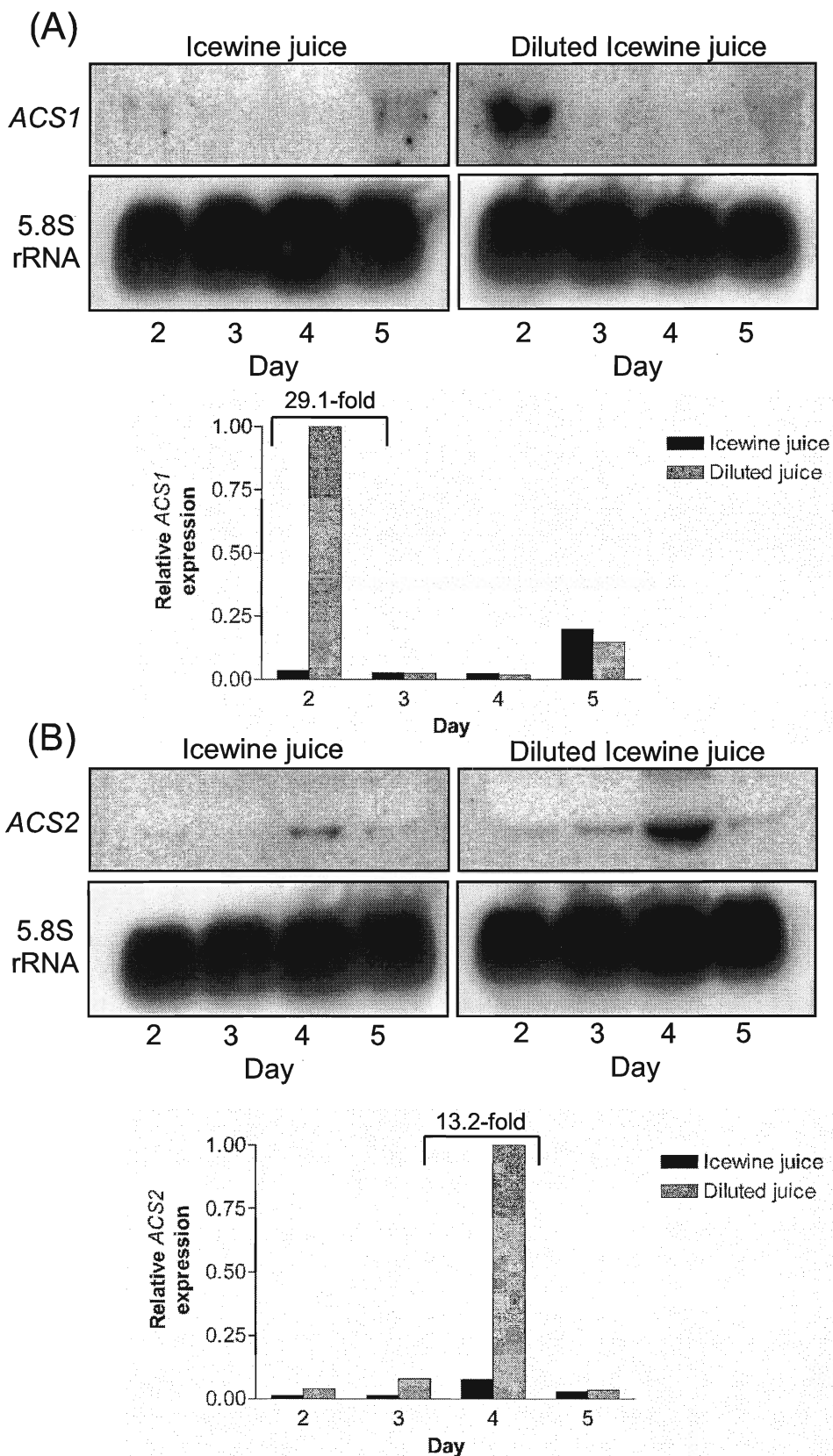


Figure 4.5.4: Expression of *ACS1* and *ACS2* during fermentation. Expression profiles of *ACS1* (A) and *ACS2* (B) from yeast fermenting Icewine juice and diluted Icewine juice using Northern analysis. 5.8S ribosomal RNA was used as a loading control. Relative gene expression between fermentation conditions is illustrated in the bar graphs.

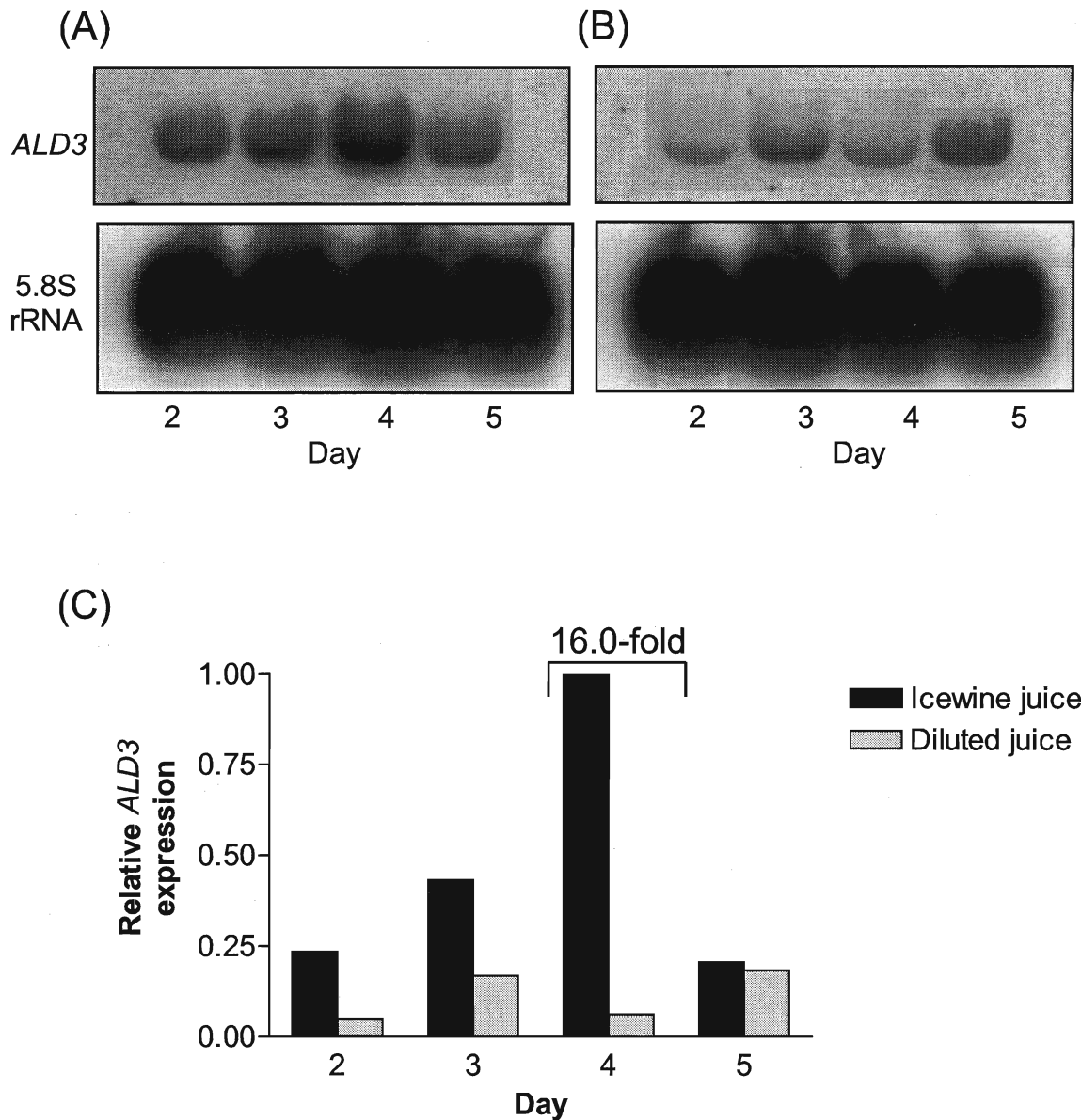


Figure 4.5.5: Expression of *ALD3* during fermentation. The expression profiles of *ALD3* from yeast fermenting Icewine juice (A) and diluted Icewine juice (B) using Northern analysis. 5.8S ribosomal RNA was used as a loading control. (C) Comparison of the relative levels of *ALD3* expression between fermentation conditions.

4.5.2. Discussion- Acetic acid Metabolism

During alcoholic fermentation, acetic acid is synthesized in *S. cerevisiae* via the cytosolic pyruvate dehydrogenase (PDH) bypass which reroutes the pyruvate produced during glycolysis away from the PDH reaction that converts pyruvate to acetyl-CoA and towards aldehyde dehydrogenase forming acetic acid. Acetic acid synthesis has been suggested as a mechanism through which yeast can reestablish cytosolic redox balance by regenerating NADH from the NAD^+ produced during glycerol biosynthesis in response to salt induced hyperosmotic stress (Blomberg and Adler, 1989; Wang *et al.*, 1998; Navarro-Avino *et al.*, 1999). Of the five genes encoding for aldehyde dehydrogenases identified in *S. cerevisiae*, upregulation of *ALD3* and *ALD6* has been associated with increased acetic acid production in osmotically stressed cells (Blomberg and Adler, 1989; Miralles and Serrano, 1995; Navarro-Avino *et al.*, 1999; Norbeck and Blomberg, 2000). However, both isoforms differ in their cofactor requirement with Ald3p being specific for NAD^+ and Ald6p being specific for NADP^+ . Due to the differences in cofactor requirement between these two *ALD* isoforms, the function of the NADP^+ -dependent dehydrogenase in maintaining redox balance in yeast under hyperosmotic stress is unclear as they lack transhydrogenase activity to convert reducing equivalents between the NAD^+/NADH and $\text{NADP}^+/\text{NADPH}$ cofactors systems (van Dijken and Scheffers, 1986; Meaden *et al.*, 1997; Navarro-Avino *et al.*, 1999). The expression of several aldehyde dehydrogenase isoforms have been implicated in acetic acid production during table wine fermentation. However, the results from the microarray data from this study in addition to the Northern analysis results from previous studies by Pigeau and Inglis (2005 and 2007) shows that it is likely that the upregulation of the NAD^+ -dependent *ALD3*

isoform contributes to the increased levels of acetic acid in Icewine. Northern analysis verified the microarray results showing that *ALD3* was also expressed throughout both fermentation conditions, however its expression was significantly greater in the Icewine juice condition with the largest fold difference in expression occurring 96 h into the Icewine juice fermentation. Also from the microarray results, the other aldehyde dehydrogenase isoforms were observed to be expressed but the expression level of these genes was not significantly different between fermentation conditions after day two.

Expression of the mitochondrial aldehyde dehydrogenase isoforms encoded by *ALD4* and *ALD5* together with the cytosolic NADP⁺-dependent isoform encoded by *ALD6* were found to contribute to acetic acid production in wine yeast strains fermenting 20% (w/v) glucose media similar to conditions found in a table wine fermentation (Remize *et al.*, 2000; Saint-Prix *et al.*, 2004). Based on these results, these three Aldp isoforms along with the cytosolic NAD⁺-dependent isoform Ald2p, likely contributed to acetic acid production in yeast during diluted Icewine juice fermentation, but their roles in the increased acetic acid synthesis and redox balancing in Icewine juice fermenting cells may be only minor. However, Erasmus *et al.* (2003) reported a significant upregulation of *ALD2*, *ALD3*, *ALD4* and *ALD6* expression in wine yeast following a 2 h exposure to 40% (w/v) sugar stress using microarray analysis. Their results may reflect a transient transcriptional response of *ALD* expression upon exposure to high sugar levels. The results from our study reflect the adapted response of wine yeast to prolonged exposure to high osmotic stress during the early stages of Icewine juice fermentation. Similar to the results reported by Pigeau and Inglis in 2005 and 2007, our microarray data indicate that the aldehyde dehydrogenases encoded by *ALD2*, 4, 5 and 6 were not

differentially expressed between the high and low sugar stress fermentation conditions, thereby indicating that these isoforms likely do not contribute to the elevated levels acetic acid produced during Icewine fermentation. From the microarray data, both mitochondrial isoforms (encoded by *ALD4* and *ALD5*) along with the cytosolic NADP⁺-dependent isoform (*ALD6*) were actually found to be significantly downregulated in Icewine juice fermenting cells on day two which would further limit their contribution to acetic acid production.

Elevated acetic acid production by Icewine juice fermenting yeast may also be implicated in the inactivation of the plasma membrane aquaglyceroporin Fps1p. In industrial *S. cerevisiae* strains used in the food industry, resistance and adaptation to weak acid stress involves the loss of Fps1p that passively facilitates the uptake of acetic acid (reviewed in Mollapour *et al.*, 2008). Acetic acid is small enough to gain entry into yeast cells through Fps1p by passive diffusion but when glucose repressed cells are confronted with high levels of acetic acid, the sudden influx of this weak acid through Fps1p can transiently activate Hog1p via the Sln1p branch of the HOG pathway (Mollapour and Piper, 2007; reviewed in Mollapour *et al.*, 2008). Hog1p in turn, phosphorylates Fps1p, therefore targeting Fps1p for endocytosis and degradation within the vacuole (Mollapour and Piper, 2007; reviewed in Mollapour *et al.*, 2008). High levels of acetic acid can initiate apoptosis in yeast cells so removing Fps1p may be part of an essential process to protect cells against apoptotic cell death when challenged with acetic acid (Giannattasio *et al.*, 2005; reviewed in Mollapour *et al.*, 2008). It is unknown if these events occur in yeast fermenting Icewine juice. By day two, the Icewine fermenting cells have already produced a considerable amount of acetic acid compared to the dilute juice

cells. Additionally from the microarray data, *FPS1* was found to be expressed by yeast under both fermentation conditions on day two but this channel rapidly closes upon exposure to hyperosmotic stress to control the amount of glycerol retained by the cell (Támas *et al.*, 1999). Activity of Fps1p in yeast during fermentation may be subject to a number of controls including high osmotic pressure and the presence of acetic acid but further research is necessary to clearly define its role during Icewine fermentation.

Since acetic acid is an intermediate of the PDH bypass in *S. cerevisiae*, acetic acid can be converted to acetyl-CoA via an ATP-dependent reaction catalyzed by an acetyl-CoA synthetase encoded by two isoforms, *ACS1* and *ACS2*. The bypass may have an important role in contributing to the cytosolic pool of acetyl-CoA which is required for lipid, sterol and amino acid synthesis (reviewed in Pronk *et al.*, 1996). *ACS1* encodes for the mitochondrial isozyme while *ACS2* encodes for the cytosolic acetyl-CoA synthetase. Although *ACS1* expression is required for respiratory growth on nonfermentable carbon sources and is glucose repressed (van den Berg *et al.*, 1996), it was strongly expressed in yeast on day two of the dilute juice fermentation. An increase in mitochondrial acetyl-CoA reserves may correspond to an increase in its consumption in other metabolic pathways. Genes associated with the TCA cycle were more highly expressed in dilute juice fermenting cells and an increase in *ACS1* expression may be required to replenish acetyl-CoA. The fatty acid β -oxidation pathway results in the formation of acetyl-CoA so an increase in the expression of genes associated with this pathway in dilute juice fermenting yeast cells may reflect a need for an increase in acetyl-CoA production. Since the rate of yeast biomass accumulation was faster in the dilute juice condition compared to the rate observed in the Icewine juice condition (see Section 4.2.1.), increased

production acetyl-CoA in yeast during dilute juice fermentation may have been required to support increased cell growth and reproduction.

However, the isoenzyme encoded by *ACS2* is required for fermentative growth on glucose (van den Berg and Steensma, 1995). *ACS2* was significantly downregulated up to day four of the Icewine fermentation. A decreased demand for cytosolic acetyl-CoA during Icewine fermentation corresponds to a decrease in the expression of genes involved in other metabolic pathways requiring the input of acetyl units such as lipid and ergosterol synthesis. Since yeast cells accumulate twice the biomass when fermenting dilute juice in comparison to the fermentation of Icewine juice, these cells would have an increased metabolic need for lipid and ergosterol synthesis. A wine yeast strain overexpressing *ACS2* did not result in reduced acetic acid levels (Remize *et al.*, 2000), however a reduction in acetic acid in sake yeast overexpressing *ACS2* was reported by Akamatsu *et al.* (2000). So, increased synthesis of acetyl-CoA by *Acs2p* from the acetic acid produced by yeast fermenting dilute juice may have resulted in the lower acetic acid levels found in this juice. But, the combination of increased *ALD3* expression together with decreased *ACS1/2* expression in osmotically stressed yeast cells during Icewine fermentation may have ultimately contributed to the elevated acetic acid levels measured in the final wine. Acetyl-CoA can also be converted back to acetic acid in a reaction catalyzed by an acetyl-CoA hydrolase encoded by *ACH1*. *ACH1* was more highly expressed in yeast fermenting dilute juice up to day four. The expression of this gene may be involved in regulating the cytosolic acetyl-CoA pools.

S. cerevisiae is able to use acetic acid as sole carbon source under aerobic conditions and the plasma membrane protein encoded by *ADY2* was identified as a

putative acetate transporter (Paiva *et al.*, 2004). Transport analyses of a $\Delta ady2$ laboratory yeast strain demonstrated that this gene mediates acetate transport which is crucial for growth on acetic acid. *ADY2* was strongly upregulated more than ten-fold in Icewine juice fermenting cells. After day two, the extracellular acetic acid levels increased considerably in the Icewine fermentation condition corresponding to an increase in the expression of this gene. It is not known if this transporter is capable of exporting acetic acid from the cell especially in wine yeast under fermentative conditions. But, due to the low concentration of acetic acid in the initial Icewine juice, the concentration gradient across the yeast plasma membrane would favour acetic acid export.

4.6. ADDITIONAL TRANSCRIPTOME RESPONSES DURING ICEWINE FERMENTATION

4.6.1. Results - additional transcriptome responses

4.6.1.1. Stress Responsive Genes

Significant fold-changes in the expression of a number of stress response genes in wine yeast were observed throughout the first five days of Icewine juice fermentation, particularly genes associated with heat shock, osmotic and oxidative stress (Table 4.6.1). Upon exposure, the expression of several yeast genes is regulated by stress with a large proportion of them responding in a related manner to many types of stresses (Causton *et al.*, 2001; Gasch *et al.*, 2000). There were six genes encoding for heat shock proteins that were found to be differentially expressed between fermentation conditions. Typically, heat shock proteins function as molecular chaperones that facilitate protein refolding and/or maintain protein conformation in yeast subject to heat-induced stress (Dawes, 2004). Of these genes, *HSP12* was strongly expressed throughout the Icewine juice fermentation with the greatest fold difference (8.9-fold) occurring on day five of the

Table 4.6.1: Expression profile of the genes involved in responding to stress.

| ORF | Gene | Description | Fold Change* | | | |
|-----------------------|--------|--|--------------|-------|-------|-------|
| | | | Day 2 | Day 3 | Day 4 | Day 5 |
| Heat Shock | | | | | | |
| YOR020C | HSP10 | mitochondrial chaperonin | + | - | - | +2.4 |
| YFL014W | HSP12 | cytosolic heat shock protein | +4.7 | +3.8 | +8.5 | +8.9 |
| YBR072W | HSP26 | cytosolic heat shock protein | -3.0 | • | • | • |
| YCR021C | HSP30 | plasma membrane heat shock protein | -3.5 | • | - | • |
| YPL240C | HSP82 | ATPase activity | • | • | • | -2.9 |
| YBL075C | SSA3 | ATPase activity | - | - | -2.1 | -5.1 |
| Osmotic Stress** | | | | | | |
| YMR169C | ALD3 | aldehyde dehydrogenase | +8.4 | +9.7 | +14.6 | × |
| YDL022W | GPD1 | glycerol-3-phosphate dehydrogenase | + | +4.3 | +6.6 | × |
| YIL053W | GPP1 | glycerol-3-phosphatase | +3.1 | +8.7 | +12.3 | +3.5 |
| YER062C | GPP2 | glycerol-3-phosphatase | +3.0 | +9.3 | +14.0 | +4.7 |
| YPL223C | GRE1 | unknown function | +2.1 | +2.7 | +6.7 | +7.5 |
| YOL151W | GRE2 | oxidoreductase activity | × | +13.7 | +24.9 | +6.1 |
| YHR104W | GRE3 | aldo-keto reductase activity | +2.9 | +2.5 | +2.7 | • |
| YMR175W | SIP18 | phospholipid binding | • | +2.1 | +4.7 | +25.4 |
| Oxidative Stress | | | | | | |
| YGR088W | CTT1 | catalase | +3.0 | +2.0 | +3.6 | • |
| YKL026C | GPX1 | glutathione peroxidase | • | -2.3 | × | × |
| YBR244W | GPX2 | glutathione peroxidase | - | -2.1 | -2.1 | • |
| YIR037W | HYR1 | glutathione peroxidase | - | -2.2 | -2.3 | • |
| YNL036W | NCE103 | carbonate dehydratase | + | +2.8 | +5.1 | +13.8 |
| YJR104C | SOD1 | copper-zinc superoxide dismutase | -2.6 | • | • | • |
| YHR008C | SOD2 | manganese superoxide dismutase | - | -2.7 | -2.1 | • |
| YML028W | TSA1 | thioredoxin peroxidase | • | -2.9 | -2.6 | -3.0 |
| YDR453C | TSA2 | thioredoxin peroxidase | × | -3.0 | -2.3 | -2.0 |
| Response to Metal Ion | | | | | | |
| YHR053C | CUP1-1 | metallothionein; copper ion binding | • | +14.4 | +15.4 | +6.7 |
| YHR055C | CUP1-2 | metallothionein; copper ion binding | • | +13.4 | +15.3 | +6.2 |
| Response to Stimulus | | | | | | |
| YML070W | DAK1 | dihydroxyacetone kinase | • | +2.3 | +2.7 | × |
| YOL053C | DDR2 | DNA damage responsive | +2.0 | • | +2.6 | +3.6 |
| YHR044C | DOG1 | 2-deoxyglucose-6-phosphate phosphatase | +7.3 | × | × | × |
| YHR043C | DOG2 | 2-deoxyglucose-6-phosphate phosphatase | +9.2 | × | +2.4 | × |
| YML004C | GLO1 | glyoxylase | + | +2.1 | +3.2 | • |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

The level of gene expression changes in *S. cerevisiae* fermenting Icewine juice in comparison to gene expression changes in *S. cerevisiae* fermenting dilute Icewine juice are shown as either positive (+ fold upregulation) or negative (-fold downregulation) values. A plus sign (+) or minus sign (-) alone indicates significant difference in gene expression but less than a 2-fold change; • indicates no significant difference in gene expression; × indicates no expression was detected.

***GPD1*, *GPP1*, *GPP2* and *ALD3* are also included in this subcategory and further discussed in Sections 4.4 and 4.5

fermentation despite being reported as subject to glucose repression (de Groot *et al.*, 2000). The expression of the majority of heat shock encoding genes profiled using microarray analysis were found to be transiently downregulated at least three-fold on day two in the Icewine condition relative to the dilute condition (Table 4.6.1). This included *HSP26* which encodes for a small cytosolic heat shock protein and *HSP30* which encodes for protein possibly involved in H^+ homeostasis (Dawes, 2004). *HSP12* and *HSP26* expression had been previously found to be induced in laboratory yeast strains under salt-induced osmotic stress (Rep *et al.*, 2000; Yale and Bohnert, 2001). *HSP82* encodes for a plasma membrane-bound protein with ATPase activity (Richter *et al.*, 2002) and expression of this gene was observed to be nearly three-fold higher on day five of the dilute Icewine juice fermentation. *SSA3* encodes for a protein that belongs to the *HSP70* family of ATP-binding chaperones which functions to prevent the aggregation of unfolded proteins (Dawes, 2004). *SSA3* expression was induced under salt stress (Rep *et al.*, 2000) but its expression was significantly reduced more than two-fold on days four and five of the Icewine juice fermentation.

According to the GO Slim functional category of genes associated with the response to stress within the *Saccharomyces* genome database (SGD), a number of these genes are further divided into subcategories including the response to osmotic stress. The genes involved in glycerol (*GPD1*, *GPP1* and *GPP2*) and acetic acid (*ALD3*) synthesis fall into this category and their expression profiles have been previously described in Sections 4.4 and 4.5. From the microarray data, the genes associated with the osmotic response appeared to be expressed early during the Icewine juice fermentation and restricted to only a few responsive genes. The strongly osmoreponsive genes (*GRE1*,

GRE2, *GRE3* and *SIP18*) were induced in yeast on day two of the Icewine juice fermentation and maintained higher expression levels throughout the days examined in comparison to the expression levels of these genes in yeast fermenting the dilute juice (Table 4.6.1). Both *GRE1* and *SIP18* encode for hydrophilin-like proteins and expression of these genes was found to be upregulated in laboratory yeast strains under desiccation stress (Garay-Arroyo *et al.*, 2000; Singh *et al.*, 2005). The fold difference in *GRE1* (7.5-fold) and *SIP18* (25.4-fold) expression peaked on day five of the Icewine juice fermentation. Expression of all three *GRE* isoforms was found to be induced by osmotic and other stresses (Garay-Arroyo and Covarrubias, 1999). *GRE2* encodes for a protein with oxidoreductase activity while *GRE3* expression is associated with methylglyoxal metabolism (Garay-Arroyo and Covarrubias, 1999; Aguilera and Prieto, 2001). *GRE2* was more highly expressed in the Icewine juice cells than *GRE3*. *GRE2* was identified as being one of the mostly highly upregulated genes (25-fold) identified in yeast fermenting Icewine juice (Table 4.6.1).

The genes associated with responding to oxidative stress were predominantly less highly expressed during Icewine juice fermentation compared to the dilute juice fermentation by approximately two-fold on days three and four of the fermentations. This expression profile was characterized by the downregulation of genes known to be associated with reactive oxygen species defense and superoxide radical detoxification such as the superoxide dismutases (*SOD1* and *SOD2*), glutathione peroxidases (*GPX1*, *GPX2* and *HYR1*) and the thioredoxin peroxidases (*TSA1* and *TSA2*). However, cytosolic catalase T (*CTT1*) which is involved in dissipating H₂O₂ into water and oxygen (Dawes, 2004; reviewed in Hohmann, 2002) was highly expressed from days two to four of the

Icewine juice fermentation relative to the dilute juice fermentation. *CTT1* expression is regulated by the general stress transcription factors Msn2p and Msn4p (Martinez-Pastor *et al.*, 1996; reviewed in Hohmann, 2002). *NCE103* encodes for a carbonate dehydratase and expression of this gene is typically weak under anaerobic conditions (Götz *et al.*, 1999). However, *NCE103* was found to be strongly expressed on day five of the Icewine juice fermentation (Table 4.6.1).

Genes encoding for copper ion binding metallothionein proteins (*CUP1-1* and *CUP1-2*) were two of the most strongly expressed genes in yeast fermenting Icewine juice. In yeast, these proteins function as copper ion scavengers in response to metal ion stress (Dawes, 2004) but under Icewine juice fermentation conditions both isoforms were expressed more than ten-fold higher on days three and four compared to the dilute fermentation (Table 4.6.1).

The response to stimulus is a broad category encompassing a subset of genes expressed under a variety of stress conditions also including osmotic and oxidative stresses. Of these genes, two of them associated with glycerol degradation (*DAK1*) and methylglyoxal metabolism (*GLO1*) exhibited differential expression between fermentation conditions (Table 4.6.1). *DAK1* encodes for a dihydroxyacetone kinase that functions to phosphorylate dihydroxyacetone to form DHAP at the expense of one ATP molecule (reviewed in Blomberg, 2000). This gene was expressed just over two-fold greater on day two in the Icewine juice when compared to the dilute juice fermentation. *GLO1* encodes an enzyme (glyoxylase I) involved in the first step in the methylglyoxal degradation pathway that ultimately forms lactic acid in the presence of glutathione (Inoue and Kimura, 1995). The difference in *GLO1* expression was at least two-fold

higher on days three and four of Icewine juice fermentation relative to the dilute juice fermentation. Two genes encoding for putative 2-deoxyglucose-6-phosphate phosphatases (*DOG1* and *DOG2*) were strongly but transiently expressed on day two of the Icewine fermentation (Table 4.6.1).

4.6.1.2. Transport Responses

Several genes associated with transport of small molecules and ions in *S. cerevisiae* were differentially expressed between fermentation conditions. Transport protein encoding genes were significantly up- or downregulated up to day four of the Icewine juice fermentation and by day five, no expression of these genes could be detected in either fermentation condition (Table 4.6.2).

The *ENA* gene family encodes for plasma membrane Na^+ -ATPase pumps mediating the active efflux of sodium from the cytoplasm (reviewed in Hohmann, 2002). *ENA1* expression was found to be induced in response to salt stress in laboratory yeast (Posas *et al.*, 2000) however three *ENA* isoforms were strongly expressed in yeast fermenting Icewine juice compared to yeast fermenting dilute juice. Also, genes involved in iron transport (*FIT1*, *FIT2* and *FTR1*) were relatively more highly expressed early on in the Icewine juice fermentation. Three genes encoding for organic acid transporters in yeast (*DIC1*, *JEN1* and *SFC1*) were found to be less highly expressed on day two of the Icewine juice fermentation. However, the difference in *SFC1* (mitochondrial succinate-fumarate antiporter) expression was more than ten-fold on day five of the fermentations (Table 4.6.2). The role of *SFC1* in yeast during Icewine fermentation is unclear since its expression is required for aerobic growth on ethanol and acetate (Fernández *et al.*, 1994; Palmieri *et al.*, 1997).

Table 4.6.2: Expression profile of the genes involved in transport.

| ORF | Gene | Description | Fold Change* | | | |
|------------------------|------|--|--------------|-------|-------|-------|
| | | | Day 2 | Day 3 | Day 4 | Day 5 |
| Ion Transport | | | | | | |
| YFL050C | ALR2 | magnesium cation transporter | × | + | +3.0 | × |
| YDR040C | ENA1 | sodium ion transporter | • | +3.5 | +5.0 | × |
| YDR039C | ENA2 | sodium ion transporter | -2.0 | +3.9 | +4.7 | × |
| YDR038C | ENA5 | sodium ion transporter | • | +3.0 | × | × |
| YDR534C | FIT1 | siderophore transporter | +2.3 | × | × | × |
| YOR382W | FIT2 | siderophore transporter | +6.5 | × | × | × |
| YER145C | FTR1 | high affinity iron permease | • | +4.7 | × | × |
| YJR077C | MIR1 | phosphate transporter | + | -2.2 | -2.8 | • |
| YER053C | PIC2 | phosphate transporter | + | - | -2.3 | -2.0 |
| YNL055C | POR1 | phosphate transporter | • | -4.6 | -3.4 | -2.6 |
| YGL255W | ZRT1 | high affinity zinc ion transporter | -6.9 | × | × | × |
| YLR130C | ZRT2 | low affinity zinc ion transporter | -3.0 | × | × | × |
| Organic Acid Transport | | | | | | |
| YLR348C | DIC1 | dicarboxylic acid transporter | -2.3 | × | × | × |
| YKL217W | JEN1 | lactate and pyuvate permease | -2.0 | • | -2.4 | × |
| YJR095W | SFC1 | mitochondrial succinate-fumarate transporter | -3.4 | + | • | +11.4 |

*Significantly different as determined by a paired t-test ($P \leq 0.05$)

The level of gene expression changes in *S. cerevisiae* fermenting Icewine juice in comparison to gene expression changes in *S. cerevisiae* fermenting dilute Icewine juice are shown as either positive (+ fold upregulation) or negative (-fold downregulation) values. A plus sign (+) or minus sign (-) alone indicates significant difference in gene expression but less than a 2-fold change; • indicates no significant difference in gene expression; × indicates no expression was detected.

4.6.2. Discussion – additional transcriptome responses

The high osmotic pressure yeast encountered during Icewine fermentation elicited the upregulation of a number of genes categorized as stress responsive according to the Gene Ontology (GO) Slim term annotations of the *Saccharomyces* genome database (SGD) (Ashburner *et al.*, 2000; Dwight *et al.*, 2002). The expression of many of these genes encode for proteins and enzymes that are primarily involved with the protection of yeast cells against osmotic and oxidative damage upon exposure to salt, oxidizing agents and other environmental stimuli (Gasch *et al.*, 2000; Rep *et al.*, 2000; Causton *et al.*, 2001). In order to select and/or design yeast strains better suited to Icewine fermentations, identifying which genes that are characteristically expressed during the adaptive stress response of commercial wine yeast under extended hyperosmotic stress throughout Icewine fermentation is of great interest.

Of the genes annotated by the SGD as responsive to osmotic stress, *SIP18* was one of the mostly highly expressed genes by yeast fermenting Icewine juice compared to yeast fermenting dilute juice. *SIP18* was one of four novel genes found to be inducible upon salt stress, and expression of this gene was found to be Hog1p-dependent (Miralles and Serrano, 1995). *SIP18* was also found to be upregulated four-fold in a wine yeast strain exposed to high sugar stress compared to nonstressed cells (Erasmus *et al.*, 2003). Rossignol *et al.* (2003) reported that *SIP18* was strongly expressed in a commercial wine strain throughout the fermentation of synthetic grape must containing 200 g L⁻¹ sugar with maximum *SIP18* expression occurring at the stationary growth phase. In agreement with the findings of Rossignol *et al.* (2003), the peak difference in *SIP18* expression occurred when Icewine and diluted juice fermenting cells entered early stationary phase.

The function of *SIP18* (salt induced protein) in response to hyperosmotic stress has not been fully elucidated but there is evidence that this gene putatively encodes for a hydrophilin (Garay-Arroyo *et al.*, 2000).

The hydrophilins are part of a broad class of proteins which include plant late embryogenesis abundant (*LEA*) proteins that are implicated in providing desiccation tolerance to plants under water-limiting conditions (Baker *et al.*, 1988; Garay-Arroyo *et al.*, 2000). *LEA*-like proteins have been identified in fungi and they appear to be involved in pathways that respond to osmotic stress and desiccation (Garay-Arroyo *et al.*, 2000; Singh *et al.*, 2005). These hydrophilic proteins are structurally characterized by high glycine contents and contain many charged amino acid residues (Bray, 1997; Garay-Arroyo *et al.*, 2000). Microarray analysis of the response of a laboratory strain of *S. cerevisiae* to desiccation and subsequent rehydration under minimal glucose conditions showed that *SIP18* was found to be strongly expressed in conjunction with another gene, *GRE1* (Singh *et al.*, 2005). In *S. cerevisiae*, 12 hydrophilin-encoding genes have been identified within the genome (Garay-Arroyo *et al.*, 2000). Of these, three (*SIP18*, *HSP12* and *GRE1*) have been previously reported to be induced in yeast exposed to osmotic stress (Miralles and Serrano, 1995; Varela *et al.*, 1992; Garay-Arroyo and Covarrubias, 1999).

Both *HSP12* and *GRE1* were also observed to be upregulated in yeast under high sugar stress and during alcoholic fermentation (Erasmus *et al.*, 2003; Rossignol *et al.*, 2003). *HSP12* is a member of the heat shock protein family (Dawes, 2004) and is typically upregulated under in yeast under a variety of stress conditions including heat shock and salt stress (Gasch *et al.*, 2000; Rep *et al.*, 2000; Causton *et al.*, 2001).

Although *HSP12* expression was shown to be subject to glucose repression (de Groot *et al.*, 2000), our microarray data indicated that the fold-change in expression of both *HSP12* and *GRE1* was higher during the first five days of Icewine juice fermentation compared to dilute juice fermentation. Both *HSP12* and *GRE1* were additionally observed to be upregulated in wine yeast during high sugar stress and alcoholic fermentation (Erasmus *et al.*, 2003; Rossignol *et al.*, 2003). Singh *et al.* (2005) proposed that in desiccated yeast that have not biosynthesized or accumulated sufficient amounts of glycerol or trehalose required for survival during desiccation, the Sip18p protein could function as a compatible solute. It has been clearly shown in previous studies that yeast synthesize and accumulate glycerol as a compatible osmolyte in response to osmotic stress (reviewed in Nevoigt and Stahl, 1997). In our present study, yeast fermenting Icewine juice also produced glycerol in response to high concentrations of solutes present in the juice (Section 4.3.1). Although yeast cells experience loss of cytosolic water and reduced turgor pressure upon exposure to osmotic stress (Blomberg and Adler, 1992; Varela and Mager, 1996; Blomberg and Adler, 2002; Billi and Potts, 2002), it is unknown if hydrophilins have a role in protecting wine yeast cells against hyperosmotic stress during Icewine fermentation.

As previously stated, *GRE1* was highly expressed in yeast fermenting Icewine juice but the microarray data also showed that two other genes, *GRE2* and *GRE3* followed the same trend. *GRE1*, *GRE2* and *GRE3* were all reported to be responsive to sorbitol-induced osmotic stress and other stresses including ionic, oxidative and heat stress (Garay-Arroyo and Covarrubias, 1999). *GRE1* and *GRE2* were also upregulated in a laboratory yeast strain in response to desiccation but the expression of these genes

declined upon yeast rehydration (Singh *et al.*, 2005). Of the *GRE* genes, only *GRE2* was significantly upregulated in wine yeast under high sugar stress 2 h post inoculation (Erasmus *et al.*, 2003). Expression of all three *GRE* genes are under control of the HOG-pathway but activation *GRE1* and *GRE3* upon stress is also mediated by the general stress response STRE-transcription factors Msn2p and Msn4p (Garay-Arroyo and Covarrubias, 1999). Amino acid sequence analysis revealed that *GRE2* encodes for a protein with similarity to a plant NADP⁺-dependent dihydroflavonol-4-reductase while *GRE3* encodes for a NADPH-dependent keto-aldose reductase highly related to fungal xylose-reductases (Garay-Arroyo and Covarrubias, 1999). There is evidence that the aldose reductase encoded by *GRE3* has a key role in the methylglyoxal detoxification pathway in yeast (Aguilera and Prieto, 2001). Both *GRE2* and *GRE3* encode for putative enzymes involved in redox reactive pathways so they could possibly participate in maintaining intracellular redox balance in the NADP⁺/NADPH cofactor system if an imbalance exists during osmotic stress. However, the function of the *GRE* genes in wine yeast fermenting Icewine juice is unknown. *ALD6*, which encodes for a NADP⁺-dependent aldehyde dehydrogenase, was found to be expressed by yeast fermenting both Icewine and diluted Icewine juice (see Table 4.5.2 in Section 4.5). Ald6p was found to be the main aldehyde dehydrogenase responsible for acetic acid production in laboratory *S. cerevisiae* strains fermenting glucose media (Remize *et al.*, 1999; Eglinton *et al.*, 2002). In addition, *ALD6* expression was found to be both upregulated under salt stress and dependent on HOG pathway activation (Akhtar *et al.*, 1997; Rep *et al.*, 2000). However, the function of Ald6p, if any, in maintaining redox balance under these hyperosmotic stress conditions during glycerol biosynthesis is not clear due to its cofactor specificity. However, Ald6p

activity in yeast may be partially responsible for the increased acetic acid produced during Icewine fermentation and may have a role in maintaining redox balance in the NADP⁺/NADPH cofactor system in order to offset the NADP⁺ generated by Gre3p during fermentation.

Expression of genes associated with responding to reactive oxygen species and superoxide radical detoxification- superoxide dismutases (*SOD1* and *SOD2*), glutathione peroxidases (*GPX1*, *GPX2* and *HYR1*) and the thioredoxin peroxidases (*TSA1* and *TSA2*)- were for the most part lower in yeast during Icewine juice fermentation compared to those fermenting dilute juice. The expression of many genes encoding proteins and enzymes involved in oxidative damage defense and in redox metabolism is activated under a variety of stress conditions (Causton *et al.*, 2001; reviewed in Hohmann, 2002). One of which, cytosolic catalase T (*CTT1*), was upregulated during in yeast fermenting high sugar grape juice (Erasmus *et al.*, 2003). *CTT1* is involved in H₂O₂ dissipation (Dawes, 2004) and this gene was also found to be upregulated during Icewine juice fermentation. Activation of *CTT1* expression is mediated by the Hog1p and the general stress response transcription factors Msn2p/Msn4p (Martinez-Pastor *et al.*, 1996; reviewed in Hohmann, 2002). Hohmann (2002) proposed that the response to oxidative stress is partially activated in yeast upon exposure to various stress conditions. When yeast are under stress, oxidative metabolism is stimulated leading to enhanced intracellular production of reactive oxygen species. However, it is unknown if expression of oxidative defense genes in yeast cells fermenting diluted juice compared to those fermenting Icewine juice actually corresponds to an increase in the generation of reactive oxygen species under anaerobic conditions.

Genes encoding enzymes involved in methylglyoxal (MG) detoxification, *GLO1* (glyoxylase I) and *GRE3* (aldose reductase) were also more highly expressed in yeast fermenting Icewine juice. Expression of *GLO1* has been found to be induced in laboratory yeast strains under salt and sorbitol stress (Inoue *et al.*, 1998). Methylglyoxal (2-oxoaldehyde) is a highly reactive cytotoxic by-product derived from dihydroxyacetone phosphate (DHAP) following the spontaneous dephosphorylation of endiolate phosphate formed from the interconversion of the glycolytic triose phosphate intermediates DHAP and glyceraldehyde-3-phosphate (Inoue *et al.*, 1998; Aguilera and Prieto, 2001; Margarida *et al.*, 2001; Aguilera *et al.*, 2005). Up to 0.3% of glucose consumed in exponentially growing yeast cells is converted to methylglyoxal (Margarida *et al.*, 2001). There are two routes of methylglyoxal detoxification in yeast. The first route involves Glo1p which requires glutathione to convert methylglyoxal to S-D-lactoylglutathione and Glo2p (glyoxylase II) catalyzes the hydrolysis of the thioester forming D-lactate and regenerating glutathione (Margarida *et al.*, 2001). *GLO2* was not found to be expressed by yeast under either fermentation condition. In the alternate pathway, Gre3p catalyzes the NADPH-dependent conversion of methylglyoxal to 1,2-propanediol (Aguilera *et al.*, 2005). Intracellular methylglyoxal levels have been shown to transiently increase upon brief hyperosmotic shock (Aguilera and Prieto, 2001). *GLO1*, but not *GRE3*, was upregulated in wine yeast under high sugar stress (Erasmus *et al.*, 2003). It is not known if yeast cells accumulate methylglyoxal when exposed to prolonged hyperosmotic stress such as during Icewine fermentation. Enhanced methylglyoxal formation should result if a proportional increase in the rate of carbon flux through glycolysis also occurs (Aguilera and Prieto, 2001). But, we observed that sugar uptake and the expression of glycolytic

genes are greatly reduced in yeast during Icewine fermentation compared to dilute juice fermentation. Upregulation of *GLO1* and *GRE3* in these cells may only be in response to the higher osmotic pressure placed on yeast in Icewine juice compared to those in diluted juice and not due to an accumulation of methylglyoxal. However, methylglyoxal accumulation in Icewine fermenting yeast could occur as a consequence of an increased demand for DHAP required for glycerol synthesis. This demand would favour the redirection of triose phosphates formed during glycolysis towards DHAP production and away from glyceraldehyde-3-phosphate thereby increasing the chances of methylglyoxal formation.

Of the several genes associated with transport of small molecules and ions in *S. cerevisiae* that were differentially expressed between fermentation conditions, the genes encoding for magnesium, sodium and iron transporters were upregulated in Icewine juice fermenting yeast. The uptake or export of ions plays an important role during osmoadaptation in all types of organisms including yeast (reviewed in Hohmann, 2002). The *ENA* gene family encodes for plasma membrane Na^+ -ATPase pumps mediating the efflux of sodium from the cytoplasm (reviewed in Hohmann, 2002). *ENA1* expression was found to be induced in response to salt stress in laboratory yeast (Posas *et al.*, 2000) and expression during osmotic stress is Hog1p-dependent (reviewed in Hohmann, 2002). Two *ENA* isoforms (*ENA2* and *ENA5*) were upregulated in sugar-stressed wine yeast (Erasmus *et al.*, 2003) however our microarray results revealed that three *ENA* isoforms were strongly upregulated in yeast fermenting Icewine juice. Additionally, genes encoding for the copper ion binding metallothionein proteins (*CUP1-1* and *CUP1-2*) were observed to be strongly expressed during Icewine fermentation. The

metallothioneins can function as antioxidants in yeast under oxidative stress conditions in addition to acting as scavengers for metal ions thereby protecting cells against copper toxicity (Dawes, 2004). It has been previously shown that Icewine juice contains concentrated solutes in addition to sugar that increases stress on wine yeast fermenting this juice (Pigeau and Inglis, 2005). Although the concentrations of salt and metal ions present in Icewine juice has not been measured to date, their presence could potentially intensify the overall hyperosmotic stress wine yeast face during Icewine fermentation and initiate other stress responses if they disrupt intracellular ion homeostasis during fermentation.

5. OVERALL DISCUSSION AND FUTURE DIRECTIONS

The successful commercial development of Canadian Icewine into a unique specialty dessert wine has garnered Canada international recognition as a world leader in Icewine production. Although the high concentrations of naturally concentrated solutes in the juice ultimately contribute to the intense flavours and aromas characteristic of this wine, the hyperosmotic nature of the juice places commercial wine yeast under considerable stress as they ferment the juice into wine (Kontkanen *et al.*, 2004; Pigeau and Inglis, 2005, Pigeau *et al.*, 2007). In this study, the high osmotic stress experienced by the commercial wine yeast *Saccharomyces cerevisiae* K1-V1116 throughout the course of Icewine fermentation elicited changes in cell growth and metabolism which correlated to several fermentation difficulties including reduced biomass accumulation and fermentation rate, reduced nutrient uptake and increased production of potentially undesirable metabolites like acetic acid that could ultimately affect the quality of the finished wine. With the use of microarray analysis, we further bridged the gap between the genome-wide transcriptional response of wine yeast as they adapt to prolonged exposure to high osmotic stress to changes within metabolic pathways associated with sugar and nitrogen and the production of yeast metabolites throughout Icewine juice fermentation. Typically throughout table wine fermentations, yeast growth is restricted by carbon and nitrogen source availability, but more frequently by nitrogen source limitation. Despite the fact that the Icewine juice contained approximately twice the amount of both sugar and nitrogen in comparison to the diluted Icewine juice, only about half the amount of available reducing sugars and amino acid nitrogen was ultimately consumed by the Icewine juice fermenting yeast. Based on these results, sugar and

nitrogen availability was not a contributing factor to the impaired cell growth exhibited by Icewine juice fermenting cells. However, the osmotic stress due to the overall abundance of concentrated sugar and nitrogen in addition to other solutes in the juice hindered the ability of the wine yeast to ferment this juice. In 2005, a study by Pigeau and Inglis demonstrated that yeast fermenting chaptalized diluted Icewine juice, which contained equal amounts of fermentable sugars as the undiluted Icewine juice but half the amounts of all other soluble solids, were able to generate more biomass and consume more sugar than Icewine juice fermenting cells. This further indicated that in addition to sugar, other solutes concentrated in the juice affects the overall fermentative capacity of wine yeast. Using microarray and Northern analysis, we also identified the expression of a glycerol/H⁺ symporter (*STL1*) which was highly upregulated in yeast fermenting Icewine juice along with the expression of two acetyl-CoA synthetase isoforms (*ACSI* and 2) which were greatly downregulated. The expression patterns of these genes potentially contribute to decreased cell viability and increased acetic acid production demonstrated by yeast fermenting Icewine juice.

The concentrated solutes in the Icewine juice increased the osmotic stress placed on yeast cells was manifested in a longer lag phase and fewer actively budding cells at the start of the fermentation. Yeast fermenting Icewine juice did not reach the same cell concentration or accumulate the same amount of biomass compared to cells fermenting diluted Icewine juice. Since there were high concentrations of residual sugar and nitrogenous compounds measured in Icewine, reduced biomass formation was not due to limited nutrient availability. The gene profiling data clearly indicated that genes associated with sugar and amino acid uptake and metabolism were predominantly

downregulated during the exponential growth phase of Icewine fermenting yeast correlating to an overall decrease in the rate of sugar and nitrogen utilization compared to yeast fermenting diluted juice. Despite the high concentrations of fermentable sugars present in the juice, many of the genes associated with sugar metabolism that were not highly expressed in yeast under the Icewine condition were involved in glycolysis and the TCA cycle. Decreased or lack of expression of the genes associated with ATP-generating metabolic pathways could contribute to impaired cell growth by limiting the amount of energy produced during Icewine fermentation that is normally required for optimal cell division and growth under anaerobic conditions. It also appears that the high osmotic stress yeast encounter during Icewine fermentation impedes amino acid uptake. In the dilute juice condition, over 90% of assimilable amino acids and ammonia were taken up after five days of the fermentation compared to only 28% in the Icewine juice condition. Reduced amino acid uptake coupled with a decrease in the expression of amino acid biosynthetic genes may also further impair cell growth and contribute nitrogen deficiencies that typically lead to sluggish fermentation rates (Monteiro and Bisson, 1991a).

Reduced biomass formation during Icewine fermentation may also be a consequence of diverting sugar away from cell growth and towards glycerol and acetic acid production as part of the yeast stress response to prolonged exposure to high osmotic conditions. The microarray results showed that the genes within the glycerol and acetic acid metabolic pathways were significantly upregulated in yeast fermenting Icewine juice and an increase in the expression of these genes corresponded to elevated levels of both metabolites found in the wine. Wine yeast face both anaerobic and osmotic stress during

Icewine fermentation and either of these factors may control the transcriptional response of genes associated with glycerol synthesis. Glycerol production by osmotically stressed yeast has been correlated to an increased expression of the NAD⁺-dependent glycerol-3-phosphate dehydrogenase (*GPD1*) in order to balance the external osmotic pressure (Larsson *et al.*, 1993; Albertyn *et al.*, 1994a; Blomberg, 2000). *GPD1* was found to upregulated during Icewine fermentation however *GPD2* was not expressed after day three by yeast under both fermentation conditions. *GPD2* isoform has been reported to serve a role in maintaining redox balance under anaerobic conditions by oxidizing excess NADH generated during biomass formation (van Dijken and Scheffers, 1986; Ansell *et al.*, 1997). In agreement with the results of Pigeau and Inglis (2005 and 2007) *GPD2*, unlike *GPD1*, does not appear to contribute to the elevated glycerol found in the wine. Both isoforms of glycerol-3-phosphatase (*GPP1* and *GPP2*) were more strongly expressed during Icewine fermentation. *GPP1* and *GPP2* are both induced by osmotic stress in a HOG pathway dependent manner (Norbeck *et al.*, 1996; Rep *et al.*, 2000; Pålman *et al.*, 2001) but *GPP2* expression was found to be more responsive to salt stress (Hirayama *et al.*, 1995) whereas *GPP1* expression was found to be transiently upregulated under anaerobic conditions (Pålman *et al.*, 2001). Although the rate-limiting step in glycerol synthesis is catalyzed by glycerol-3-phosphate dehydrogenase (Larsson *et al.*, 1993; Eriksson *et al.*, 1995; Remize *et al.*, 2001), both isoforms were highly expressed in yeast in the presence of high concentrations of fermentable sugars, and the increased expression appears to be required for glycerol synthesis during Icewine fermentation.

Active glycerol uptake via a glycerol/H⁺ symporter (Stl1p) may be occurring in *S. cerevisiae* during Icewine juice fermentation. *STL1* was one of the most strongly expressed genes identified in yeast fermenting Icewine juice. *STL1* expression has been previously described as salt-induced and glucose repressed in laboratory yeast strains (Rep *et al.*, 2000; Ferreira *et al.*, 2005). However, results from this thesis study showed *STL1* was one of the most highly upregulated genes in wine yeast. Similar results were found when wine yeast were briefly exposed to high sugar stress (Erasmus *et al.*, 2003). Using both microarray and Northern analysis in our study, an expression profile comparison of this gene throughout the Icewine juice and dilute juice fermentations resulted in a relative 25-fold difference in *STL1* expression between juice conditions on day five of the fermentations.

The use of Stl1p by yeast for glycerol uptake during Icewine juice fermentation may contribute to limited cell growth by dissipating the plasma membrane proton gradient in an energy dependent manner. Protons that enter the cell must be excreted to prevent acidification of the cytoplasm and cell death but proton export requires the activity of several plasma membrane ATPases (reviewed in Boulton *et al.*, 1998). This extra energy expenditure to maintain the proton gradient may also further disadvantage yeast during Icewine fermentation. Reduced expression of ATPase encoding genes like *HSP82* may decrease the ability of yeast cells to regulate intracellular proton concentrations during Icewine fermentation. Disruption in the proton motive force necessary to import amino acids through the dissipation of the plasma membrane proton gradient may also be potentially responsible for reduced amino acid uptake observed in Icewine juice fermenting cells.

Although expression of *STL1* is HOG pathway dependent (reviewed in Hohmann, 2002) and it appears that *STL1* expression is induced upon prolonged exposure to high sugar stress during Icewine fermentation, upregulation of *STL1* in yeast does not necessarily mean that its protein product is functional during Icewine fermentation. Future studies involving [¹⁴C]glycerol transport assays in wine yeast fermenting Icewine and diluted Icewine juice in the presence and absence of a chemical protonophore will help to determine if yeast cells are actively taking up glycerol during Icewine fermentation.

Additionally from the microarray data, upregulation of the aldehyde dehydrogenase isoform *ALD3* appears to have a key role in acetic acid formation and maintaining redox balance during glycerol synthesis in yeast during Icewine fermentation. The production of acetic acid by Ald3p can resupply the cytosolic NADH required to support increased glycerol production in Icewine fermenting yeast cells in response to the high extracellular osmotic pressure. Erasmus *et al.* (2003) reported the upregulation of *ALD2*, *ALD3*, *ALD4* and *ALD6* in wine yeast briefly exposed to high sugar stress. However, our microarray results were in agreement with the findings by Pigeau and Inglis (2005 and 2007) whereby the upregulation of *ALD3* during Icewine fermentation in conjunction with a lack of differential expression of the other aldehyde dehydrogenase isoforms (*ALD2*, 4, 5 and 6) between fermentation conditions corresponded to a considerable fold increase in acetic acid production by Icewine fermenting yeast. Expression of the mitochondrial aldehyde dehydrogenase isoforms *ALD4* and *ALD5* together with the cytosolic NADP⁺-dependent isoform *ALD6* were found to contribute to acetic acid production in wine yeast strains fermenting 20% (w/v)

glucose media similar to conditions found in a table wine fermentation (Remize *et al.*, 2000; Saint-Prix *et al.*, 2004). HOG-dependent upregulation of *ALD3* and *ALD6* has been associated with increased acetic acid production in osmotically stressed cells (Blomberg and Adler, 1989; Miralles and Serrano, 1995; Navarro-Avino *et al.*, 1999; Norbeck and Blomberg, 2000). However *ALD3* encodes for a cytosolic NAD⁺-dependent aldehyde dehydrogenase (Navarro-Avino *et al.*, 1999) while *ALD6* encodes for a cytosolic NADP⁺-dependent isoform (Meaden *et al.*, 1997) thus the redox balancing role of Ald6p during Icewine fermentation is not clear since yeast lack a transhydrogenase to convert between reducing equivalents (van Dijken and Scheffers, 1986). Even though we have transcriptional evidence of *ALD3* contributing to the elevated acetic acid levels generated by yeast fermenting Icewine juice, enzyme assays would clarify which cytosolic aldehyde dehydrogenase contributes to acetic acid production.

Acetic acid can be the substrate of acetyl-CoA synthetase which catalyzes the formation of acetyl-CoA from acetate and coenzyme A. Two acetyl-CoA synthetase isoenzymes exist in *S. cerevisiae* and the cytosolic isoenzyme encoded by *ACS2* is necessary for growth on glucose by yeast under anaerobic conditions (van den Berg and Steensma 1995) while *ACS1* is the glucose repressible mitochondrial isoform required for respiratory growth on non-fermentable carbon sources (Kratzer and Schüller, 1995; Van den Berg *et al.*, 1996). A cytosolic supply of acetyl-CoA is required for lipid, sterol and amino acid biosynthesis such that yeast can utilize the cytosolic pyruvate dehydrogenase bypass to generate cytosolic acetyl-CoA (reviewed in Pronk *et al.* 1996). Acetic acid is an intermediate of the bypass in *S. cerevisiae* and it can be converted to acetyl-CoA via an ATP-dependent reaction catalyzed by acetyl-CoA synthetase. The bypass may have an

important role in contributing to the cytosolic pool of acetyl-CoA which is required for lipid, sterol and amino acid synthesis (reviewed in Pronk *et al.*, 1996).

The microarray expression profiles of the acetyl-CoA synthetases which were successfully verified by Northern analysis showed that both *ACS* isoforms were strongly downregulated in yeast cells fermenting Icewine juice compared to their diluted juice fermenting counterparts. Downregulation of *ACS2* during Icewine fermentation corresponded to the downregulation of genes involved in other metabolic pathways requiring the input of acetyl units such as lipid, ergosterol and amino acid synthesis. Yeast cells fermenting dilute juice accumulate twice the biomass than those fermenting Icewine juice therefore demonstrating that less osmotically stressed cells have a metabolic need for increased various biosynthetic reactions and depend on replenishing the pool of available cytosolic acetyl-CoA. The combination of increased *ALD3* expression together with the decrease in *ACS1/2* expression in osmotically stressed yeast cells during Icewine fermentation may ultimately contribute to the elevated acetic acid levels measured in the wine. A reduction in acetyl-CoA formation from acetate by *ACS2* may be a contributing factor to the elevated acetic acid levels measured in Icewine. A reduction in acetic acid in sake yeast overexpressing *ACS2* was reported by Akamatsu *et al.* (2000) but a wine yeast strain also overexpressing *ACS2* did not result in reduced acetic acid levels (Remize *et al.*, 2000). It is unclear if increased expression of *ACS2* during the diluted juice fermentation contributes to increased acetic acid catabolism and therefore lower levels of acetic acid released into the wine. It is also not known if increasing *ACS2* expression in a wine yeast strain during Icewine fermentation would promote increased acetic acid catabolism. Since *ACS1* is expressed during respiratory

growth and is glucose repressed (van den Berg *et al.* 1996) it is unlikely that it would have an influence on increased acetic acid production during Icewine fermentation.

The gene profiling data also revealed the upregulation of a number of stress response genes including those involved in redox reactive metabolic pathways (*GRE1*, 2 and 3) and those related to oxidative stress (*CTT1*). The expression of these genes have been found to be upregulated under numerous stress conditions (Gasch *et al.*, 2000; Causton *et al.*, 2001; Rossignol *et al.*, 2003; Erasmus *et al.*, 2003), their importance to yeast under Icewine fermentations conditions clearly needs to be further investigated. A large-scale transcriptomic study by Marks *et al.* (2008) revealed that over 200 genes were highly induced by a wine yeast strain throughout the fermentation of grape juice and were categorized into their own group termed the fermentation stress response (FSR). The expression of a number of these genes affected by the stresses imposed on yeast cells during fermentation were associated with transport, response to stress but the vast majority of them are of unknown function and still have not been characterized. The microarray data generated in our study can be further mined to identify additional genes that may be uniquely up- or downregulated in response to Icewine fermentation and specifically those used to cope with high sugar stress conditions. Ando *et al.* (2006) used a collection of deletion mutants to identify genes required for tolerance to high sucrose stress in Baker's yeast. Of the 273 osmosensitive deletion mutants, 269 showed cross-sensitivities to high concentrations of sodium chloride and sorbitol. However, four genes associated with purine biosynthesis and intracellular ATP accumulation were found to be necessary for sucrose stress tolerance. Thus, the stress response of wine yeast to concentrated solutes present in Icewine juice may differ from the responses observed

with other osmotic stresses, especially since the wine yeast are also present in an anaerobic environment. These characteristic responses could possibly involve the activation of Icewine-specific genes or entire metabolic pathways not identified to date. It would also be of importance to identify the transcriptional responses of genes that are expressed constitutively or not affected during Icewine fermentation in addition to the expression of genes required for adaptation to the changing physiological and metabolic conditions faced by yeast cells during alcoholic fermentation. This would facilitate a greater understanding of the overall response of *S. cerevisiae* to Icewine fermentation thereby improving our ability to select for or design strains better suited to these fermentations.

6. CONCLUSIONS

With the use of microarray analysis, this thesis project has demonstrated that the hyperosmotic stress experienced by the commercial wine yeast *Saccharomyces cerevisiae* K1-V1116 during the first five days of Icewine fermentation compared to table wine fermentation evoked transcriptional changes on a genomic level. These differences in gene expression ultimately lead to differences in yeast cell growth and metabolism. During Icewine fermentation, yeast were observed to experience several fermentation difficulties including decreased biomass accumulation, decreased nutrient uptake and increased production of metabolites like glycerol and acetic acid. Despite the fact that the Icewine juice contained approximately twice the amount of both sugar and nitrogen in comparison to the diluted Icewine juice, only half the amount of available reducing sugars and amino acid nitrogen was utilized by the Icewine juice fermenting yeast. Based on these results, sugar and nitrogen availability was not a contributing factor to the

impaired cell growth exhibited by Icewine juice fermenting cells. However, the osmotic stress due to the overall abundance of concentrated sugar and nitrogen in addition to other solutes in the juice hindered the ability of the wine yeast to ferment this juice. This study also identified the expression of a glycerol/H⁺ symporter (*STL1*) which was one of the most highly expressed genes in yeast fermenting Icewine juice along with the expression of two acetyl-CoA synthetase isoforms (*ACS1* and 2) which were greatly downregulated. The expression patterns of these genes potentially contributed to decreased cell viability and increased acetic acid production demonstrated by yeast fermenting Icewine juice. The results presented in this study has further bridged the gap between the transcriptional response of wine yeast as they adapt to prolonged exposure to hyperosmotic stress to changes within metabolic pathways associated with sugar and nitrogen and the production of yeast metabolites throughout Icewine juice fermentation thereby improving our ability to select for yeast strains better suited to these fermentations.

LITERATURE CITED

Aguilera, J. and Prieto, J.A. (2001) The *Saccharomyces cerevisiae* aldose reductase is implied in the metabolism of methylglyoxal in response to stress conditions. *Curr Genet* **39**:273-283.

Aguilera, J., Rodríguez-Vargas, S. and Prieto, J.A. (2005) The HOG MAP kinase pathway is required for the induction of methylglyoxal-responsive genes and determines methylglyoxal resistance in *Saccharomyces cerevisiae*. *Mol Microbiol* **56**:228-239.

Akamatsu, S., Kamiya, H., Yamashita, N., Motoyoshi, T., Goto-Yamamoto, N., Ishikawa, T., Okazaki, N. and Nishimura, A. (2000) Effects of aldehyde dehydrogenase and acetyl-CoA synthetase on acetate formation in sake mash. *J Biosci Bioeng* **90**:555-560.

Akhtar, N., Blomberg, A. and Adler, L. (1997) Osmoregulation and protein expression in a *pbs2Δ* mutant of *Saccharomyces cerevisiae* during adaptation to hypersaline stress. *FEBS Lett* **403**:173-180.

Albertyn, J., Hohmann, S. and Prior, B.A. (1994a) Characterization of the osmotic-stress response in *Saccharomyces cerevisiae*: osmotic stress and glucose repression regulate glycerol-3-phosphate dehydrogenase independently. *Curr Genet* **25**:12-18.

Albertyn, J., Hohmann, S., Thevelein, J.M. and Prior, B.A. (1994b) *GPD1*, which encodes glycerol-3-phosphate dehydrogenase, is essential for growth under osmotic stress *Saccharomyces cerevisiae*, and its expression is regulated by the high-osmolarity glycerol response pathway. *Mol Cell Biol* **14**:4135-4144.

Alepuz, P.M., Jovanovic, A., Reiser, V. and Ammerer, G. (2001) Stress-Induced MAP Kinase Hog1 Is Part of Transcription Activation Complexes. *Mol Cell* **7**:767-777.

Andre, B., Hein, C., Grenson, M. and Jauniaux, J.C. (1993) Cloning and expression of the *UGA4* gene coding for the inducible GABA-specific transport protein of *Saccharomyces cerevisiae*. *Mol Gen Genet* **237**:17-25.

Ando, A., Tanaka, F., Murata, Y., Takagi, H. and Shima, J. (2006) Identification and classification of genes required for tolerance to high-sucrose stress by genome-wide screening of *Saccharomyces cerevisiae*. *FEMS Yeast Res* **6**:249-267.

Ansell, R., Granath, K., Hohmann, S., Thevelein, J.M. and Adler, L. (1997) The two isoenzymes for yeast NAD^+ -dependent glycerol 3-phosphate dehydrogenase encoded by *GPD1* and *GPD2* have distinct roles in osmoadaptation and redox regulation. *EMBO J* **16**:2179-2187.

- Ashburner, M., Ball, C.A., Blake, J.A., Botstein, D., Butler, H., Cherry, J. M., Davis, A.P., Dolinski, K., Dwight, S.S., Eppig, J.T., Harris, M.A., Hill, D.P., Issel-Tarver, L., Kasarskis, A., Lewis, S., Matese, J.C., Richardson, J.E., Ringwald, M., Rubin, G.M. and Sherlock, G. (2000) Gene Ontology: tool for the unification of biology. *Nature Genet* **25**:25-29.
- Backhus, L.E., DeRisi, J., Brown, P.O. and Bisson, L.F. (2001) Functional genomic analysis of a commercial wine strain of *Saccharomyces cerevisiae* under differing nitrogen conditions. *FEMS Yeast Res* **1**:111-125.
- Baker, J., Steele, C. and Dure, L. (1988) Sequence and characterization of 6 *Lea* proteins and their genes from cotton. *Plant Mol Biol* **11**:277-291.
- Beltran, G., Novo, M., Rozès, M., Mas, A. and Guillamón, J.M. (2004) Nitrogen catabolite repression in *Saccharomyces cerevisiae* during wine fermentation. *FEMS Yeast Res* **4**:625-632.
- Bidenne, C., Blondin, B., Dequin, S. and Vezinhet, E. (1992) Analysis of the chromosomal DNA polymorphism of wine strains of *Saccharomyces cerevisiae*. *Curr Genet* **22**:1-7
- Berthels, N.J., Otero, R.R.C., Bauer, F.F., Thevelein, J.M. and Pretorius, I.S. (2004) Discrepancy in glucose and fructose utilisation during fermentation by *Saccharomyces cerevisiae* wine yeast strains. *FEMS Yeast Res* **4**:683-689.
- Billi, D. and Potts, M. (2002) Life and death of dried prokaryotes. *Res Microbiol* **153**:7-12.
- Bisson, L.F. (1993) Yeasts - Metabolism of sugars. In *Wine Microbiology and Biotechnology* Fleet, G.H. (Ed.) pp. 55-75. Switzerland: Harwood Academic Publishers.
- Bisson, L.F. and Fraenkel, D.G. (1983) Involvement of kinases in glucose and fructose uptake by *Saccharomyces cerevisiae*. *Proc Natl Acad Sci USA* **80**:1730-1734.
- Bisson, L.F. and Fraenkel, D.G. (1984) Expression of Kinase-Dependent Glucose Uptake in *Saccharomyces cerevisiae*. *J Bacteriol* **159**:1013-1017.
- Blomberg, A. (2000) Metabolic surprises in *Saccharomyces cerevisiae* during adaptation to saline conditions: questions, some answers and a model. *FEMS Microbiol Lett* **182**:1-8.
- Blomberg, A. and Adler, L. (1989) Roles of glycerol and glycerol-3-phosphate dehydrogenase (NAD⁺) in acquired osmotolerance of *Saccharomyces cerevisiae*. *J Bacteriol* **171**:1087-1092.

- Blomberg, A. and Adler, L. (1992) Physiology of osmotolerance in fungi. *Adv Microb Physiol* **33**:145–212
- Blomberg, A., Larsson, C. and Gustafsson, L. (1988) Microcalorimetric monitoring of growth of *Saccharomyces cerevisiae*: Osmotolerance in relation to physiological state. *J Bacteriol* **170**:4562-4568.
- Boulton, R.B., Singleton, V.L., Bisson, L.F. and Kunkee, R.E. (1998) *Principles and practices of winemaking*. Maryland: Aspen Publishers, Inc.
- Bray, E.A. (1997) Plant responses to water deficit. *Trends Plant Sci* **2**:48-54.
- Brewster, J.L., de Valoir, T., Dwyer, N.D., Winter, E. and Gustin, M.C. (1993) An osmosensing signal transduction pathway in yeast. *Science* **259**: 1760-1763.
- Camarasa, C., Grivet, J. and Dequin, S. (2003) Investigation by ^{13}C -NMR and tricarboxylic acid (TCA) deletion mutant analysis of pathways for succinate formation in *Saccharomyces cerevisiae* during anaerobic fermentation. *Microbiology* **149**:2669-2678.
- Causton, H.C., Ren, B., Koh, S.S., Harbison, C.T., Kanin, E., Jennings, E.G., Lee, T.I., True, H.L., Lander, E.S. and Young, R.A. (2001) Remodeling of yeast genome expression in response to environmental changes. *Mol Biol Cell* **12**:323–337.
- Cavalieri, D., Townsend, J.P. and Hartl, D.L. (2000) Manifold anomalies in gene expression in a vineyard isolate of *Saccharomyces cerevisiae* revealed by DNA microarray analysis. *Proc Natl Acad Sci USA* **97**:12369-12374.
- Cliff, M.A. and Pickering, G.J. (2006) Determination of odour detection thresholds for acetic acid and ethyl acetate in Ice wine. *J Wine Res* **17**:45-52.
- Coleman, S.T., Fang, T.K., Rovinsky, S.A., Turano, F.J. and Moye-Rowley, W.S. (2001) Expression of a glutamate decarboxylase homologue is required for normal oxidative stress tolerance in *Saccharomyces cerevisiae*. *J Biol Chem* **276**:244-250.
- Cooper, T.G. (1982) Nitrogen metabolism in *Saccharomyces cerevisiae*. In *The Molecular Biology of the Yeast Saccharomyces: Metabolism and Gene Expression*. Strathern, J.N., Jones, E.W. and Broach, J.R. (Eds.) pp 39-99. New York: Cold Spring Harbor Laboratory.
- Cordier, H., Mendes, F., Vasconcelos, I. and François, J.M. (2007) A metabolic and genomic study of engineered *Saccharomyces cerevisiae* strains for high glycerol production. *Metab Eng* **9**:364-378.
- Dawes, I.W. (2004) Stress responses. In *The Metabolism and Molecular Physiology of Saccharomyces cerevisiae* Dickinson, J.R. and Schweizer, M. (Eds.) pp. 376-413. New York: CRC Press.

- de Groot, E., Bebelman, J., Mager, W.H. and Planta, R. J. (2000) Very low amounts of glucose cause repression of the stress-responsive gene *HSP12* in *Saccharomyces cerevisiae*. *Microbiology* **146**:367-375.
- Denis, C.L., Ferguson, J. and Young, E.T. (1983) mRNA levels for the fermentative alcohol dehydrogenase of *Saccharomyces cerevisiae* decrease upon growth on a nonfermentable carbon source. *J Biol Chem* **258**:1165-1171.
- DeRisi, J.L., Iyer, V.R. and Brown, P.O. (1997) Exploring the metabolic and genetic control of gene expression on a genomic scale. *Science* **278**:680–686.
- Dickinson, J.R., Salgado, L. and Hewlins, M. (2003) The catabolism of amino acids to long chain and complex alcohols in *Saccharomyces cerevisiae*. *J Biol Chem* **278**:8028-8034.
- Dickinson, J.R. (2004) Nitrogen metabolism. In *The Metabolism and Molecular Physiology of Saccharomyces cerevisiae* Dickinson, J.R. and Schweizer, M. (Eds.) pp. 104-116. New York: CRC Press.
- DeLuna, A., Avendano, A., Riego, L. and González, A. (2001) NADP-glutamate dehydrogenase isoenzymes of *Saccharomyces cerevisiae*. *J Biol Chem* **276**:43775-43783.
- Dukes, B.C. and Butzke, C.E. (1998) Rapid determination of primary amino acids in grape juice using an *o*-Phthaldialdehyde/N-Acetyl-L-Cysteine spectrophotometric assay. *Am J Enol Vitic* **49**:125-134.
- Dwight, S.S., Harris, M.A., Dolinski, K., Ball, C.A., Binkley, G., Christie, K.R., Fisk, D.G., Issel-Tarver, L., Schroeder, M., Sherlock, G., Sethuraman, A., Weng, S., Botstein, D. and Cherry, M. (2002) *Saccharomyces* Genome Database (SGD) provides secondary gene annotation using the Gene Ontology (GO). *Nucleic Acids Res* **30**:69-72
- Eglinton, J.M., Heinrich, A.J., Pollnitz, A.P., Langridge, P., Henschke, P.A. and de Barros Lopes, M. (2002) Decreasing acetic acid accumulation by a glycerol overproducing strain of *Saccharomyces cerevisiae* by deleting the *ALD6* aldehyde dehydrogenase gene. *Yeast* **19**:295-301.
- Erasmus, D.J., van der Merwe, G.K. and van Vuuren, H.J.J. (2003) Genome-wide expression analyses: Metabolic adaptation of *Saccharomyces cerevisiae* to high sugar stress. *FEMS Yeast Res* **3**:375-399.
- Eriksson, P., André, L., Ansell, R., Blomberg, A. and Adler, L. (1995) Cloning and characterization of *GPD2*, a second gene encoding *sn*-glycerol 3-phosphate dehydrogenase (NAD⁺) in *Saccharomyces cerevisiae*, and its comparison with *GPD1*. *Mol Microbiol* **17**:95-107.

- Eriksson, P., Alipour, H., Adler, L. and Blomberg, A. (2000) Rap1p-binding sites in the *Saccharomyces cerevisiae* GPD1 promoter are involved in its response to NaCl. *J Biol Chem* **275**:29368–29376.
- Esteve-Zarzoso, B., Belloc h, C., Uruburu, F. and Querol, A. (1999) Identification of yeasts by RFLP analysis of the 5.8S rRNA gene and the two ribosomal internal transcribed spacers. *Int J Syst Bacteriol* **49**: 329–337.
- Fernández, M., Fernández, E. and Rodicio, R. (1994) *ACR1*, a gene encoding a protein related to mitochondrial carriers, is essential for acetyl-CoA synthetase activity in *Saccharomyces cerevisiae*. *Mol Gen Genet* **242**:727-735.
- Ferreira, C., van Voorst, F., Martins, A., Neves, L., Oliveira, R., Kielland-Brandt, M.C., Lucas, C. and Brandt, A. (2005) A member of the sugar transporter family, Stl1p is the glycerol/H⁺ symporter in *Saccharomyces cerevisiae*. *Mol Biol Cell* **16**:2068-2076.
- Ferrigno, P., Posas, F., Koepp, D., Saito, H. and Silver, P.A. (1998) Regulated nucleo/cytoplasmic exchange of HOG1 MAPK requires the importin β homologs NMD5 and XPO1. *EMBO J* **17**:5606–5614.
- Fleet, G.H. and Heard, G.M (1993) Yeasts- Growth during Fermentation. In *Wine Microbiology and Biotechnology* Fleet, G.H. (Ed.) pp. 27-54. Switzerland: Harwood Academic Publishers.
- Flikweert, M.T., van der Zanden, L., Janssen, W.M., Steensma, H.Y., van Dijken, J.P. and Pronk, J.T. (1996) Pyruvate decarboxylase: an indispensable enzyme for growth of *Saccharomyces cerevisiae* on glucose. *Yeast* **12**:247-257.
- Gancedo, J.M. (1998) Yeast carbon catabolite repression. *Microbiol Mol Biol Rev* **62**:334-361.
- Garay-Arroyo, A. and Covarrubias, A.A. (1999) Three genes whose expression is induced by stress in *Saccharomyces cerevisiae*. *Yeast* **15**:879-892.
- Garay-Arroyo, A., Colmenero-Flores, J.M., Garcíarrubio, A. and Covarrubias, A.A. (2000) Highly hydrophilic proteins in prokaryotes and eukaryotes are common during conditions of water deficit. *J Biol Chem* **275**:5668-5674.
- Gasch, A.P., Spellman, P.T., Kao, C.M., Carmel-Harel, O., Eisen, M.B., Storz, G., Botstein, D. and Brown, P.O. (2000) Genomic Expression Programs in the Response of Yeast Cells to Environmental Changes. *Mol Biol Cell* **11**:4241–4257.
- Giannattasio, S., Guaragnella, N., Corte-Real, M., Passarellac, S. and Marra, E. (2005) Acid stress adaptation protects *Saccharomyces cerevisiae* from acetic acid-induced programmed cell death. *Gene* **354**:94-98.

Godard, P., Urrestarazu, A., Vissers, S., Kontos, K., Bontempi, G., van Helden, J. and André, B. (2007) Effect of 21 different nitrogen sources on global gene expression in the yeast *Saccharomyces cerevisiae*. *Mol Cell Biol* **27**:3065-3086.

Görner, W., Durchschlag, E., Martinez-Pastor, M.T., Estruch, F., Ammerer, G., Hamilton, B., Ruis, H. and Schüller, C. (1998) Nuclear localization of the C₂H₂ zinc finger protein Msn2p is regulated by stress and protein kinase A activity. *Gene Dev* **12**:586-597.

Götz, R., Gnann, A. and Zimmerman, F.K. (1999) Deletion of the carbonic anhydrase-like gene *NCE103* of the yeast *Saccharomyces cerevisiae* causes an oxygen-sensitive growth defect. *Yeast* **15**:855-864.

Gustin, M.C., Albertyn, J., Alexander, M. and Davenport, K. (1998) MAP Kinase Pathways in the yeast *Saccharomyces cerevisiae*. *Microbiol Mol Biol Rev* **62**:1264-1300.

Hauser, N.C., Fellenberg, K., Gil, R., Bastuck, S., Hoheisel, J.D. and Pérez-Ortín, J.E. (2001) Whole genome analysis of a wine yeast strain. *Comp Funct Genom* **2**:69-79.

Herrero, P., Galindez, J., Ruiz, N., Martinez-Campa, C. and Moreno, F. (1995) Transcriptional regulation of the *Saccharomyces cerevisiae* *HXK1*, *HXK2* and *GLK1* genes. *Yeast* **11**:137-144.

Hiltunen, J.K., Mursula, A.M., Rottensteiner, H., Wierenga, R.K., Kastaniotis, A.J. and Gurvitz, A. (2003) The biochemistry of peroxisomal β -oxidation in the yeast *Saccharomyces cerevisiae*. *FEMS Microbiol Rev* **27**:35-64.

Hirayama, T., Maeda, T., Saito, H. and Shinozaki, K. (1995) Cloning and characterization of seven cDNAs for hyperosmolarity-responsive (*HOR*) genes of *Saccharomyces cerevisiae*. *Mol Gen Genet* **249**:127-138.

Hohmann, S. (1991) *PCD6*, a weakly expressed pyruvate decarboxylase gene from yeast, is activated when fused spontaneously under the control of the *PDC1* promoter. *Curr Genet* **20**:373-378.

Hohmann, S. and Cederberg, H. (1990) Autoregulation may control the expression of yeast pyruvate decarboxylase structural genes *PDC1* and *PDC5*. *Eur J Biochem* **188**:615-621.

Hohmann, S. (2002) Osmotic stress signaling and osmoadaptation in yeasts. *Microbiol Mol Biol Rev* **66**:300-372.

Holst, B., Lunde, C., Lages, F., Oliveira, R., Lucas, C. and Kielland-Brandt, M.C. (2000) *GUP1* and its close homologue *GUP2*, encoding multimembrane-spanning proteins involved in active glycerol uptake in *Saccharomyces cerevisiae*. *Mol Microbiol* **37**:108-124.

- Horák, J. (1986) Amino acid transport in eucaryotic microorganisms. *Biochim Biophys Acta* **864**:223-256.
- Inoue, Y. and Kimura, A. (1995) Methylglyoxal and regulation of its metabolism in microorganisms. *Adv Microb Physiol* **37**:177-227.
- Inoue, Y., Tsujimoto, Y. and Kimura, A. (1998) Expression of the glyoxalase I gene of *Saccharomyces cerevisiae* is regulated by high osmolarity glycerol mitogen-activated protein kinase pathway in osmotic stress response. *J Biol Chem* **273**:2977-2983.
- Jacoby, T., Flanagan, H., Faykin, A., Seto, A.G., Mattison, C. and Ota, I. (1997) Two protein-tyrosine phosphatases inactivate the osmotic stress response pathway in yeast by targeting the mitogen-activated protein kinase, Hog1. *J Biol Chem* **272**:17749–17755.
- Jansen, G., Buhring, F., Hollenberg, C.P. and Ramezani Rad, M. (2001) Mutations in the SAM domain of STE50 differentially influence the MAP kinase-mediated pathways for mating, filamentous growth and osmotolerance in *Saccharomyces cerevisiae*. *Mol Genet Genom* **265**:102–117.
- Kinclova-Zimmermannova, O. and Sychrova, H. (2006) Functional study of the Nha1p C-terminus: involvement in cell response to changes in external osmolarity. *Curr Genet* **49**:229-236.
- Kluba, R.M., Mattick, L.R. and Hackler, L.R. (1978) Changes in the free and total amino acid composition of several *Vitis labruscana* grape varieties during maturation. *Am J Enol Vitic* **29**:102-111.
- Kontkanen, D., Inglis, D.L., Pickering, G.J. and Reynolds, A. (2004) Effect of yeast inoculation rate, acclimatization and nutrient addition on Icewine fermentation. *Am J Enol Vitic* **55**:363-370.
- Kratzer, S. and Schüller, H. (1995) Carbon source-dependent regulation of the acetyl-coenzyme A synthetase-encoding gene *ACS1* from *Saccharomyces cerevisiae*. *Gene* **161**:75-79.
- Krukeberg, A.L. and Dickinson, J.R. (2004) Carbon metabolism. In *The Metabolism and Molecular Physiology of Saccharomyces cerevisiae* Dickinson, J.R. and Schweizer, M. (Eds.) pp. 42-103. New York: CRC Press.
- Lages, F. and Lucas, C. (1997) Contribution to the physiological characterization of glycerol active uptake in *Saccharomyces cerevisiae*. *Biochim Biophys Acta* **1322**:8-18.
- Larsson, K., Ansell, R., Eriksson, P. and Adler, L. (1993) A gene encoding sn-glycerol 3-phosphate dehydrogenase (NAD⁺) complements an osmosensitive mutant of *Saccharomyces cerevisiae*. *Mol Microbiol* **10**:1101-1111.

- Latterich, M. and Watson, M.D. (1993) Evidence for a dual osmoregulatory mechanism in the yeast *Saccharomyces cerevisiae*. *Biochem Biophys Res Commun* **191**:1111–1117.
- Lee, F., Lin, L. and Smith, J.A. (1996) Acetyl-CoA hydrolase involved in acetate utilization in *Saccharomyces cerevisiae*. *Biochim Biophys Acta* **1297**:99-104.
- Lee, J., Lee, B., Shin, D., Kwak, S., Bahk, J.D., Lim, C.O. and Yun, D. (2002) Carnitine uptake by *AGP2* in yeast *Saccharomyces cerevisiae* is dependent on Hog1 MAP kinase pathway *Mol Cells* **13**:407-412.
- Luyten, K., Albertyn, J., Skibbe, W.F., Prior, B.A., Ramos, J., Thevelein, J.M. and Hohmann, S. (1995) Fps1, a yeast member of the MIP family of channel proteins, is a facilitator for glycerol uptake and efflux and is inactive under osmotic stress. *EMBO J* **14**:1360-1371.
- Luyten, K., Riou, C. and Blondin, B. (2002) The hexose transporters of *Saccharomyces cerevisiae* play different roles during enological fermentation. *Yeast* **19**:713-726.
- Maeda, T., Tsai, A.Y.M. and Saito, H. (1993) Mutations in a protein tyrosine phosphatase gene (PTP2) and a protein serine/threonine phosphatase gene (PTC1) cause a synthetic growth defect in *Saccharomyces cerevisiae*. *Mol Cell Biol* **13**:5408–5417.
- Maeda, T., Wurgler-Murphy, S.M. and Saito, H. (1994) A two-component system that regulates an osmosensing MAP kinase cascade in yeast. *Nature* **369**:242-245.
- Maeda, T., Takekawa, M. and Saito, H. (1995) Activation of yeast PBS2 MAPKK by MAPKKs or by binding of an SH3-containing osmosensor. *Science* **269**:554-558.
- Magasanik, B. and Kaiser, C.A. (2002) Nitrogen regulation in *Saccharomyces cerevisiae*. *Gene* **290**:1-18.
- Mager, W.H. and Siderius, M. (2002) Novel insights into the osmotic stress response of yeast. *FEMS Yeast Res* **2**:251-257.
- Margarida, A., Cordeiro, C., Ponces Freire, A. (2001) In situ analysis of methylglyoxal metabolism in *Saccharomyces cerevisiae*. *FEBS Lett* **499**:41-44.
- Marks, V.D., van der Merwe, G.K. and van Vuuren, H.J.J. (2003) Transcriptional profiling of wine yeast in fermenting grape juice: regulatory effect of diammonium phosphate. *FEMS Yeast Res* **3**:269-287.
- Marks, V.D., Ho Sui, S.J., Erasmus, D., van der Merwe, G.K., Brumm, J., Wasserman, W.W., Bryan, J. and van Vuuren, H.J.J. (2008) Dynamics of the yeast transcriptome during wine fermentation reveals a novel fermentation stress response. *FEMS Yeast Res* **8**:35-52.

Martinez-Pastor, M.T., Marchler, G., Schüller, C., Marchler-Bauer, A., Ruis, H. and Estruch, F. (1996) The *Saccharomyces cerevisiae* zinc finger proteins Msn2p and Msn4p are required for transcriptional induction through the stress-response element (STRE). *EMBO J* **15**:2227–2235.

Meaden, P., Dickinson, F., Mifsud, A., Tessier, W., Westwater, J., Bussey, H. and Midgley, M. (1997) The *ALD6* gene of *Saccharomyces cerevisiae* encodes a cytosolic, Mg^{2+} -activated acetaldehyde dehydrogenase. *Yeast* **13**:1319-1327.

Miralles, V.J. and Serrano, R. (1995) A genomic locus in *Saccharomyces cerevisiae* with four genes up-regulated by osmotic stress. *Mol Microbiol* **17**:653-662.

Modig, T., Granath, K., Adler, L. and Lidén, G. (2007) Anaerobic glycerol production by *Saccharomyces cerevisiae* strains under hyperosmotic stress. *Appl Microbiol and Biotechnol* **75**:289-296.

Mollapour, M. and Piper, P.W. (2007) Hog1 mitogen-activated protein kinase phosphorylation targets the yeast Fps1 aquaglycerolporin for endocytosis, thereby rendering cells resistant to acetic acid. *Mol Cell Biol* **27**:6446-6556.

Mollapour, M., Shepherd, A. and Piper, P.W. (2008) Novel stress responses facilitate *Saccharomyces cerevisiae* in the presence of the monocarboxylate preservatives. *Yeast* **25**:169-177.

Monteiro, F.F. and Bisson, L.F. (1991a) Biological assay of nitrogen content of grape juice and prediction of sluggish fermentations. *Am J Enol Vitic* **42**:47-57.

Monteiro, F.F. and Bisson, L.F. (1991b) Amino acid utilization and urea formation during vinification fermentations. *Am J Enol Vitic* **42**:199-208.

Navarro-Avino, J.P., Prasad, R., Miralles, V.J., Benito, R.M. and Serrano, R. (1999) A proposal for nomenclature of aldehyde dehydrogenases in *Saccharomyces cerevisiae* and characterization of the stress-inducible *ALD2* and *ALD3*. *Yeast* **15**:829-842.

Nelissen, B., de Wachter, R. and Goffeau, A. (1997) Classification of all putative permeases and other membrane plurispanners of the major facilitator superfamily encoded by the complete genome of *Saccharomyces cerevisiae*. *FEMS Microbiol Rev* **21**:113–134.

Neves, L., Lages, F. and Lucas, C. (2004) New insights on glycerol transport in *Saccharomyces cerevisiae*. *FEBS Lett* **565**:160-162.

Nevoigt, E. and Stahl, U. (1997) Osmoregulation and glycerol metabolism in the yeast *Saccharomyces cerevisiae*. *FEMS Microbiol Rev* **21**:231-241.

- Noble, A.C. and Bursick, G.F. (1984) The contribution of glycerol to perceived viscosity and sweetness in white wine. *Am J Enol Vitic* **35**:110-112.
- Norbeck, J. and Blomberg, A. (1998) Amino acid uptake is strongly affected during exponential growth of *Saccharomyces cerevisiae* in 0.7 M NaCl medium. *FEMS Microbiol Lett* **158**:121-126.
- Norbeck, J. and Blomberg, A. (2000) The level of cAMP-dependent protein kinase A activity strongly affects osmotolerance and osmo-instigated gene expression changes in *Saccharomyces cerevisiae*. *Yeast* **16**:121-137.
- Norbeck, J., Pählman, A., Akhtar, N., Blomberg, A. and Adler, L. (1996) Purification and characterization of two isoenzymes of DL-glycerol-3-phosphatase from *Saccharomyces cerevisiae*. *J Biol Chem* **271**:13875-13881.
- Nurgel, C., Pickering, G.J. and Inglis, D.L. (2004) Sensory and chemical characteristics of Canadian ice wines. *J Sci Food Agr* **84**:1675-1684.
- Nurgel, C. and Pickering, G.J. (2005) Contribution of glycerol, ethanol and sugar to the perception of viscosity and density elicited by model white wines. *J Texture Stud* **36**:303-323.
- Oliveira, R., Lages, F., Silva-Graça, M. and Lucas, C. (2003) Fps1p channel is the mediator of the major part of glycerol passive diffusion in *Saccharomyces cerevisiae*: artifacts and re-definitions. *Biochim Biophys Acta* **1613**: 57-71.
- Oliveira, R. and Lucas, C. (2004) Expression studies of *GUP1* and *GUP2*, genes involved in glycerol active transport in *Saccharomyces cerevisiae*, using semi-quantitative RT-PCR. *Curr Genet* **46**:140-146.
- O'Rourke, S., Herskowitz, I. and O'Shea, E.K. (2002) Yeast go the whole HOG for the hyperosmotic stress response. *Trends Genet* **18**:405-412.
- O'Rourke, S.M. and Herskowitz, I. (2004) Unique and redundant roles for HOG MAPK pathway components as revealed by whole-genome expression analysis. *Mol Biol Cell* **15**:532-542.
- Ostrander, D.B. and Gorman, J.A. (1999) The extracellular domain of the *Saccharomyces cerevisiae* Sln1p membrane osmolarity sensor is necessary for kinase activity. *J Bacteriol* **181**:2527-2534.
- Ota, I.M. and Varshavsky, A. (1993) A yeast protein similar to bacterial two-component regulators. *Science* **262**:566-569.
- Özcan, S. and Johnston, M. (1999) Function and Regulation of Yeast Hexose Transporters. *Microbiol Mol Biol Rev* **63**:554-569.

- Påhlman, A., Granath, K., Ansell, R., Hohmann, S. and Adler, L. (2001) The yeast glycerol 3-phosphatases Gpp1p and Gpp2p are required for glycerol biosynthesis and differentially involved in the cellular responses to osmotic, anaerobic and oxidative stress. *J Biol Chem* **276**:3555-3563.
- Paiva, S., Devaux, F., Barbosa, S., Jacq, C. and Casal, M. (2004) Ady2p is essential for the acetate permease activity in the yeast *Saccharomyces cerevisiae*. *Yeast* **21**:201-210.
- Palmieri, L., Lasorsa, F.M., De Palma, A., Palmieri, F., Runswick, M.J. and Walker, J.E. (1997) Identification of the yeast *ACR1* gene product as a succinate-fumarate transporter essential for growth on ethanol or acetate. *FEBS Lett* **417**:114-118.
- Paulsen, I.T., Sliwinski, M.K., Nelissen, B., Goffeau, A. and Saier, M.H. (1998) Unified inventory of established and putative transporters encoded within the complete genome of *Saccharomyces cerevisiae*. *FEBS Lett* **430**:116-125.
- Pavlik, P., Simon, M., Schuster, T. and Ruis, H. (1993) The glycerol kinase (*GUT1*) gene of *Saccharomyces cerevisiae*: cloning and characterization. *Curr Genet* **24**:21-25.
- Perez, M., Luyten, K., Michel, R., Riou, C. and Blondin, B. (2005) Analysis of *Saccharomyces cerevisiae* hexose carrier expression during wine fermentation: both low- and high-affinity Hxt transporters are expressed. *FEMS Yeast Res* **5**:351-361.
- Perez-Torrado, R., Carrasco, P., Aranda, A., Gimeno-Alcañiza, J., Pérez-Ortina, J.E., Matallana, E. and del Olmo, M. (2002) Study of the first hours of microvinification by the use of osmotic stress-response genes as probes. *Syst Appl Microbiol* **25**:153-161.
- Pigeau, G.M. and Inglis, D.L. (2005) Upregulation of ALD3 and GPD1 in *Saccharomyces cerevisiae* during Icewine fermentation. *J Appl Microbiol* **99**:112-125.
- Pigeau, G.M. and Inglis, D.L. (2007) Response of wine yeast (*Saccharomyces cerevisiae*) aldehyde dehydrogenases to acetaldehyde stress during Icewine fermentation. *J Appl Microbiol* **103**:1576-1586.
- Pigeau, G.M., Bozza, E., Kaiser, K. and Inglis, D.L. (2007) Concentration effect of Riesling Icewine juice on yeast performance and wine acidity. *J Appl Microbiol* **103**:1691-1698.
- Posas, F., Wurgler-Murphy, S.M., Maeda, T., Witten, E.A., Thai, T.C. and Saito, H. (1996) Yeast HOG1 MAP kinase cascade is regulated by a multistep phosphorelay mechanism in the SLN1-YPD1-SSK1 "two-component" osmosensor. *Cell* **86**:865-875.
- Posas, F. and Saito, H. (1997) Osmotic activation of the HOG MAP kinase pathway via Ste11p MAPKKK: scaffold role of Pbs2p MAPKK. *Science* **276**:1702-1705.
- Posas, F. and Saito, H. (1997) Osmotic activation of the HOG MAP kinase pathway via Ste11p MAPKKK: scaffold role of Pbs2p MAPKK. *Science* **276**:1702-1705.

- Posas, F. and Saito, H. (1998) Activation of the yeast SSK2 MAP kinase kinase kinase by the SSK1 two-component response regulator. *EMBO J* **17**:1385-1394.
- Posas, F., Takekawa, M. and Saito, H. (1998) Signal transduction by MAP kinase cascades in budding yeast. *Curr Opin Microbiol* **1**:175-182.
- Posas, F., Chambers, J.R., Heyman, J.A., Hoeffler, J.P., de Nadal, E. and Ariño, J. (2000) The Transcriptional Response of Yeast to Saline Stress. *J Biol Chem* **275**:17249-17255.
- Pretorius, I.S. (2000) Tailoring wine yeast for the new millennium: novel approaches to the ancient art of winemaking. *Yeast* **16**:675-729.
- Proft, M. and Struhl, K. (2004) MAP kinase-mediated stress relief that precedes and regulates the timing of transcriptional induction. *Cell* **118**:351-361.
- Pronk, J.T., Steensma, H.Y. and van Dijken, J.P. (1996) Pyruvate metabolism in *Saccharomyces cerevisiae*. *Yeast* **12**:1607-1633.
- Radler, F. (1993) Yeasts - Metabolism of organic acids. In *Wine Microbiology and Biotechnology* Fleet, G.H. (Ed.) pp. 165-182. Switzerland: Harwood Academic Publishers.
- Raitt, D.C., Posas, F. and Saito, H. (2000) Yeast Cdc42 GTPase and Ste20 PAK-like kinase regulate Sho1-dependent activation of the Hog1 MAPK pathway. *EMBO J* **19**:4623-4631.
- Ramos, F., El Guezzer, M., Grenson, M. and Wiame, J. (1985) Mutations affecting the enzymes involved in the utilization of 4-aminobutyric acid as nitrogen source by the yeast *Saccharomyces cerevisiae*. *Eur J Biochem* **149**:401-404.
- Regenberg, B., During-Olsen, L., Kielland-Brandt, M.C. and Holmberg, S. (1999) Substrate specificity and gene expression of the amino-acid permeases in *Saccharomyces cerevisiae*. *Curr Genet* **36**:317-328.
- Reifenberger, E., Boles, E. and Ciriacy, M. (1997) Kinetic characterization of individual hexose transporters of *Saccharomyces cerevisiae* and their relation to the triggering mechanisms of glucose repression. *Eur J Biochem* **245**:324-333.
- Reiser, V., Ruis, H. and Ammerer, G. (1999) Kinase activity-dependent nuclear export opposes stress-induced nuclear accumulation and retention of Hog1 mitogen-activated protein kinase in the budding yeast *Saccharomyces cerevisiae*. *Mol Biol Cell* **10**:1147-1161.
- Reiser, V., Salah, S.M. and Ammerer, G. (2000) Polarized localization of yeast Pbs2 depends on osmotic stress, the membrane protein Sho1 and Cdc42. *Nat Cell Biol* **2**:620-627.
- Remize, F., Roustau, J.L., Sablayrolles, J.M., Barre, P. and Dequin, S. (1999) Glycerol overproduction by engineered *Saccharomyces cerevisiae* wine yeast leads to substantial changes in by-product formation and to a stimulation of fermentation rate in stationary phase. *Appl Environ Microbiol* **65**:143-149.

- Remize, F., Andrieu, E. and Dequin, S. (2000) Engineering of the pyruvate dehydrogenase bypass in *Saccharomyces cerevisiae*: role of the cytosolic Mg^{2+} and mitochondrial K^+ acetaldehyde dehydrogenases Ald6p and Ald4p in acetate formation during alcoholic fermentation. *Appl Environ Microbiol* **66**:3151-3159.
- Remize, F., Barnavon, L. and Dequin, S. (2001) Glycerol export and glycerol-3-phosphate dehydrogenase, but not glycerol phosphatase, are rate limiting for glycerol production in *Saccharomyces cerevisiae*. *Metabol Eng* **3**:301-312.
- Remize, F., Cambon, B., Barnavon, L. and Dequin, S. (2003) Glycerol formation during wine fermentation is mainly linked to Gpd1p and is only partially controlled by the HOG pathway. *Yeast* **20**:1243-1253.
- Rep, M., Albertyn, J., Thevelein, J.M., Prior, B.A. and Hohmann, S. (1999) Different signalling pathways contribute to the control of GPD1 expression by osmotic stress in *Saccharomyces cerevisiae*. *Microbiol* **145**:715-727.
- Rep, M., Krantz, M., Thevelein, J.M. and Hohmann, S. (2000) The Transcriptional Response of *Saccharomyces cerevisiae* to Osmotic Shock. *J Biol Chem* **275**: 8290-8300.
- Richter, K., Reinstein, J. and Buchner, J. (2002) N-terminal residues regulate the catalytic efficiency of the Hsp90 ATPase cycle. *J Biol Chem* **277**:44905-44910.
- Rodriguez, A., De La Cera, T., Herrero, P. and Moreno, F. (2001) The hexokinase 2 protein regulates the expression of the *GLK1*, *HXK1* and *HXK2* genes of *Saccharomyces cerevisiae*. *Biochem J* **355**:625-631.
- Rønnow, B. and Kielland-Brandt, M. (1993) *GUT2*, a gene for mitochondrial glycerol 3-phosphate dehydrogenase of *Saccharomyces cerevisiae*. *Yeast* **9**:1121-1130.
- Rossignol, T., Dulau, L., Julien, A. and Blondin, B. (2003) Genome-wide monitoring of wine yeast gene expression during alcoholic fermentation. *Yeast* **20**:1369-1385.
- Saint-Prix, F., Bönquist, L. and Dequin, S. (2004) Functional analysis of the *ALD* gene family of *Saccharomyces cerevisiae* during anaerobic growth on glucose: the $NADP^+$ -dependent Ald6p and Ald5p isoforms play a major role in acetate formation. *Microbiol* **150**:2209-2220.
- Sanders, E.M. and Ough, C.S. (1985) Determination of Free Amino Acids in Wine by HPLC. *Am J Enol Vitic* **36**:1:43-46
- Scanes, K.T., Hohmann, S. and Prior, B.A. (1998) Glycerol production by the yeast *Saccharomyces cerevisiae* and its relevance to wine: a review. *S Afr J Enol Vitic* **19**:17-24.
- Schreiner, J. (2001) Icewine: The complete story. pp. 18-25. Toronto: Warwick Publishing Inc.

- Schreve, J.L. and Garrett, J.M. (2004) Yeast Agp2p and Agp3 function as amino acid permeases in poor nutrient conditions. *Biochem Biophys Res Commun* **313**:745-751.
- Schüller, G., Brewster, J.L., Alexander, M.R., Gustin, M.C. and Ruis, H. (1994) The HOG pathway controls osmotic regulation of transcription via the stress response element (STRE) of the *Saccharomyces cerevisiae* CTT1 gene. *EMBO J* **13**:4382-4389.
- Schweizer, M. (2004) Lipids and membranes. In *The Metabolism and Molecular Physiology of Saccharomyces cerevisiae* Dickinson, J.R. and Schweizer, M. (Eds.) pp. 140-223. New York: CRC Press.
- Serrano, R. and Rodriguez-Navarro, A. (2001) Ion homeostasis during salt stress in plants. *Curr Opin Cell Biol* **13**:399-404.
- Shelp, B.J., Bown, A.W. and McLean, M.D. (1999) Metabolism and functions of gamma-aminobutyric acid. *Trends Plant Sci* **4**:446-452.
- Singh, K.K. and Norton, R.S. (1991) Metabolic changes induced during adaptation of *Saccharomyces cerevisiae* to a water stress. *Arch Microbiol* **156**:38-42.
- Singh, J., Kumar, D., Ramakrishnan, N., Singhal, V., Jervis, J., Garst, J.F., Slaughter, S.M., DeSantis, A.M., Potts, M. and Helm, R.F. (2005) Transcriptional response of *Saccharomyces cerevisiae* to desiccation and rehydration. *Appl Env Microbiol* **71**:8752-8763
- Stanbrough, M. and Magasanik, B. (1995) Transcriptional and posttranslational regulation of the general acid permease of *Saccharomyces cerevisiae*. *J Bacteriol* **177**:94-102.
- Sutherland, F.C.W., Lages, F., Lucas, C., Luyten, K., Albertyn, J., Hohmann, S., Prior, B.A. and Kilian, S.G. (1997) Characteristics of Fps1-Dependent and -Independent Glycerol Transport in *Saccharomyces cerevisiae*. *J Bacteriol* **179**:7790-7795.
- Swiegers, J.H., Dippenaar, N., Pretorius, I.S. and Bauer, F.F. (2001) Carnitine-dependent metabolic activities in *Saccharomyces cerevisiae*: three carnitine acetyltransferases are essential in a carnitine-dependent strain. *Yeast* **18**:585-595.
- Tamás, M.J., Luyten, K., Sutherland, F.C.W., Hernandez, A., Albertyn, J., Valadi, H., Li, H., Prior, B.A., Kilian, S.G., Ramos, J., Gustafsson, L., Thevelein, J.M. and Hohmann, S. (1999) Fps1p controls the accumulation and release of the compatible solute glycerol in yeast osmoregulation. *Mol Microbiol* **31**:1087-1104.
- Tamás, M.J., Rep, M., Thevelein, J.M. and Hohmann, S. (2000) Stimulation of the yeast high osmolarity glycerol (HOG) pathway: evidence for a signal generated by a change in turgor rather than by water stress. *FEBS Lett* **472**:159-165.
- ter Linde, J.J.M., Liang, H., Davis, R.W., Steensma, H.Y., van Dijken, J.P. and Pronk, J.T. (1999) Genome-wide transcriptional analysis of aerobic and anaerobic chemostat cultures of *Saccharomyces cerevisiae*. *J Bacteriol* **181**:7409-7413.

- ter Schure, E.G., van Riel, N.A.W. and Verrips, C.T. (2000) The role of ammonia metabolism in nitrogen catabolite repression in *Saccharomyces cerevisiae*. *FEMS Microbiol Rev* **24**:67-83.
- Van Aelst, L., Hohmann, S., Zimmermann, F.K., Jans, A.W.H. and Thevelein, J.M. (1991) A yeast homologue of the bovine lens fibre MIP gene family complements the growth defect of a *Saccharomyces cerevisiae* mutant on fermentable sugars but not its defect in glucose-induced RAS-mediated cAMP signaling. *EMBO J* **10**:2095-2104.
- van den Berg, M.A. and Steensma, H.Y. (1995) *ACS2*, a *Saccharomyces cerevisiae* gene encoding acetyl-coenzyme A synthetase, essential for growth on glucose. *Eur J Biochem* **231**:704-713.
- van den Berg, M.A., de Jong-Gubbels, P., Kortland, C.J., van Dijken, J.P., Pronk, J.T. and Steensma, H.Y. (1996) The two acetyl-coenzyme A synthetases of *Saccharomyces cerevisiae* differ with respect to kinetic properties and transcriptional regulation. *J Biol Chem* **271**:28953-28959.
- van Dijken, J. and Sheffers, W. (1986) Redox balances in the metabolism of sugars by yeasts. *FEMS Microbiol Rev* **32**:199-224.
- van Roermund, C.W., Elgersma, Y., Singh, N., Wanders, R.J. and Tabak, H.F. (1995) The membrane of peroxisomes in *Saccharomyces cerevisiae* is impermeable to NAD(H) and acetyl-CoA under *in vivo* conditions. *EMBO J* **14**:3480-3486.
- van Zyl, P.J., Kilian, S.G. and Prior, B.A. (1990) The role of an active mechanism in glycerol accumulation during osmoregulation by *Zygosaccharomyces rouxii*. *Appl Microbiol Biotechnol* **34**:231-235.
- Varela J.C.S., van Beekvelt, C., Planta, R.J. and Mager, W.H. (1992) Osmostress-induced changes in yeast gene expression. *Mol Microbiol* **6**:2183-2190
- Varela, J.C.S. and Mager, W.H. (1996) Response of *Saccharomyces cerevisiae* to changes in external osmolarity. *Microbiology* **142**: 721-731.
- Vintners Quality Alliance (VQA) Act (1999) Vintners Quality Alliance Act. Ontario Regulation 406/00.
- Vintners Quality Alliance Ontario (2001) Annual Report. Vintners Quality Alliance. Toronto, Ontario.
- Vintners Quality Alliance Ontario (2007) Annual Report. Vintners Quality Alliance. Toronto, Ontario.
- Visser, S., Andre, B., Muijldermans, F. and Grens, M. (1989) Positive and negative regulatory elements control the expression of the *UGA4* gene coding for the inducible 4-aminobutyric-acid-specific permease in *Saccharomyces cerevisiae*. *Eur J Biochem* **181**:357-61.

- Wang, X., Mann, C.J., Bai, Y., Ni, L. and Weiner, H. (1998) Molecular cloning, characterization, and potential roles of cytosolic and mitochondrial aldehyde dehydrogenases in ethanol metabolism in *Saccharomyces cerevisiae*. *J Bacteriol* **148**:822-830.
- Warmka, J., Hanneman, J., Lee, J., Amin, D. and Ota, I. (2001) Ptc1, a type 2C Ser/Thr phosphatase, inactivates the HOG pathway by dephosphorylating the mitogen-activated protein kinase Hog1. *Mol Cell Biol* **21**:51–60.
- West, A.H. and Stock, A.M. (2001) Histidine kinases and response regulator proteins in two-component signaling systems. *Trends Biochem Sci* **26**:369–376.
- White, W.H., Skatrud, P.L., Xue, Z. and Toyn, J.H. (2003) Specialization of function among aldehyde dehydrogenases: the *ALD2* and *ALD3* genes are required for β -alanine biosynthesis in *Saccharomyces cerevisiae*. *Genetics* **163**:69-77.
- Wieczorke, R., Krampe, S., Weierstall, T., Freidel, K., Hollenberg, C.P. and Boles, E. (1999) Concurrent knock-out of at least 20 transporter genes is required to block uptake of hexoses in *Saccharomyces cerevisiae*. *FEBS Lett* **464**:123-128.
- Wodicka, L., Dong, H., Mittmann, M., Ho, M. and Lockhart, D.J. (1997) Genome-wide expression monitoring in *Saccharomyces cerevisiae*. *Nat Biotechnol* **15**:1359-1367.
- Wurgler-Murphy, S.M., Maeda, T., Witten, E.A. and Saito, H. (1997) Regulation of the *Saccharomyces cerevisiae* Hog1 mitogen-activated protein kinase by the Ptp2 and Ptp3 protein tyrosine phosphatases. *Mol Cell Biol* **17**:1289–1297.
- Yale, J. and Bohnert, H.J. (2001) Transcript Expression in *Saccharomyces cerevisiae* at High Salinity. *J Biol Chem* **276**:15996-16007.
- Zoecklein, B.W., Fugelsang, K.C. and Gump, B.H. (1996) Laboratory Procedures. *In Wine Analysis and Production*. pp. 474–477. New York: Chapman and Hall, International Thomson Publishing.
- Zuzuarregui, A., Monteoliva, L., Gil, C. and del Olmo, M. (2006) Transcriptomic and proteomic approach for understanding the molecular basis of adaptation of *Saccharomyces cerevisiae* to wine fermentation. *Appl Environ Microbiol* **72**:836-847.

APPENDIX I RAW MICROARRAY DATA

Table 1: Raw microarray data for fermentation day two.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-------------|
| YDR384C | ATO3 | ammonium transporter | -4.7080 | -26.1 | 0.000218073 |
| YAL054C | ACS1 | acetate-CoA ligase | -4.2472 | -19.0 | 0.00136708 |
| YOR388C | FDH1 | formate dehydrogenase | -3.9233 | -15.2 | 0.00112218 |
| YPL276W | | formate dehydrogenase | -3.9048 | -15.0 | 0.0018313 |
| YIL057C | | unknown | -3.8852 | -14.8 | 0.000269461 |
| YCR010C | ADY2 | ammonium transporter | -3.7943 | -13.9 | 0.000531308 |
| YPL275W | FDH2 | formate dehydrogenase | -3.4663 | -11.1 | 0.00099206 |
| YJL152W | | | -3.1458 | -8.9 | 0.000432178 |
| YOR100C | CRC1 | carnitine:acyl carnitine antiporter | -3.1388 | -8.8 | 0.207383 |
| YPR002W | PDH1 | unknown | -3.0130 | -8.1 | 0.000701407 |
| YAR037W | | | -2.9253 | -7.6 | 0.00359492 |
| YGR236C | SPG1 | unknown | -2.8983 | -7.5 | 0.000121819 |
| YAR035W | YAT1 | carnitine O-acetyltransferase | -2.8412 | -7.2 | 0.00131289 |
| YNL117W | MLS1 | malate synthase | -2.8367 | -7.1 | 0.00327977 |
| YJL153C | INO1 | inositol-3-phosphate synthase | -2.8182 | -7.1 | 0.00723627 |
| YGL255W | ZRT1 | high affinity zinc uptake transporter | -2.7940 | -6.9 | 0.0145379 |
| YIR031C | DAL7 | malate synthase | -2.7370 | -6.7 | 0.00173296 |
| YMR107W | SPG4 | unknown | -2.7080 | -6.5 | 0.00169631 |
| YHR139C | SPS100 | unknown | -2.6807 | -6.4 | 0.000117484 |
| YCR005C | CIT2 | citrate (Si)-synthase | -2.6782 | -6.4 | 0.00406735 |
| YOR374W | ALD4 | aldehyde dehydrogenase | -2.6317 | -6.2 | 0.0107259 |
| YMR118C | | unknown | -2.5947 | -6.0 | 0.00108584 |
| YNR002C | ATO2 | ammonium transporter | -2.5923 | -6.0 | 8.52408E-05 |
| YOR108W | LEU9 | 2-isopropylmalate synthase | -2.5578 | -5.9 | 0.000410554 |
| YOR230W | WTM1 | transcription corepressor | -2.5483 | -5.8 | 0.00397148 |
| YNL104C | LEU4 | 2-isopropylmalate synthase | -2.4547 | -5.5 | 0.0014671 |
| YBL015W | ACH1 | acetyl-CoA hydrolase | -2.3808 | -5.2 | 0.00598072 |
| YKR097W | PCK1 | phosphoenolpyruvate carboxykinase | -2.3562 | -5.1 | 0.00160329 |
| YKL187C | | unknown | -2.3010 | -4.9 | 0.00213718 |
| YLR153C | ACS2 | acetate-CoA ligase | -2.2760 | -4.8 | 0.00197593 |
| YPL265W | DIP5 | amino acid transporter | -2.2202 | -4.7 | 0.00815705 |
| YER024W | YAT2 | carnitine O-acetyltransferase | -2.2080 | -4.6 | 0.00638496 |
| YLR377C | FBP1 | fructose-bisphosphatase | -2.1493 | -4.4 | 0.00168549 |
| YER073W | ALD5 | 3-chloroallyl aldehyde dehydrogenase | -2.1482 | -4.4 | 1.08085E-05 |
| YML042W | CAT2 | carnitine O-acetyltransferase | -2.1473 | -4.4 | 0.000981853 |
| YLR284C | ECI1 | dodecenoyl-CoA delta-isomerase | -2.0637 | -4.2 | 0.00124702 |
| YER065C | ICL1 | isocitrate lyase | -2.0530 | -4.1 | 0.0022822 |
| YKL182W | FAS1 | [acyl-carrier protein] S-malonyltransferase * | -2.0425 | -4.1 | 0.00282861 |
| YCR028C | FEN2 | pantothenate transporter | -2.0362 | -4.1 | 0.00153978 |
| YMR162C | DNF3 | phospholipid-translocating ATPase | -2.0222 | -4.1 | 0.00603812 |
| YGR205W | | ATP binding | -1.8952 | -3.7 | 0.00389137 |
| YPL274W | SAM3 | S-adenosylmethionine transporter | -1.8845 | -3.7 | 0.0110817 |
| YCR033W | SN11 | NAD-dependent histone deacetylase * | -1.8798 | -3.7 | 0.00195978 |
| YKL095W | YJU2 | unknown | -1.8765 | -3.7 | 0.0167637 |
| YML075C | HMG1 | hydroxymethylglutaryl-CoA reductase (NADPH) | -1.8725 | -3.7 | 0.040413 |
| YAL062W | GDH3 | glutamate dehydrogenase | -1.8695 | -3.7 | 0.00747882 |
| YCR035C | RRP43 | 3'-5'-exoribonuclease | -1.8692 | -3.7 | 0.00149969 |
| YCR050C | | | -1.8357 | -3.6 | 0.000974659 |
| YCR021C | HSP30 | unknown | -1.8237 | -3.5 | 0.00169245 |
| YOR243C | PUS7 | pseudouridine synthase | -1.8205 | -3.5 | 0.0128418 |
| YAL009W | SPO7 | unknown | -1.8122 | -3.5 | 0.00558326 |
| YNL202W | SPS19 | 2,4-dienoyl-CoA reductase (NADPH) | -1.8025 | -3.5 | 0.00130246 |
| YJR095W | SFC1 | succinate:fumarate antiporter | -1.7818 | -3.4 | 0.000114951 |
| YLR331C | | | -1.7783 | -3.4 | 0.00372839 |
| YPR006C | ICL2 | methylisocitrate lyase | -1.7735 | -3.4 | 0.000791573 |
| YBR050C | REG2 | protein phosphatase type 1 regulator | -1.7712 | -3.4 | 0.00222256 |
| YDR161W | | unknown | -1.7522 | -3.4 | 0.00168643 |
| YPL036W | PMA2 | hydrogen-exporting ATPase , phosphorylative mechanism | -1.7505 | -3.4 | 0.00103896 |
| YJL043W | | unknown | -1.7497 | -3.4 | 0.00722442 |
| YBR026C | ETR1 | enoyl-[acyl-carrier protein] reductase | -1.7342 | -3.3 | 0.00442948 |
| YDL049C | KNH1 | unknown | -1.6668 | -3.2 | 0.00182923 |
| YHR185C | PFS1 | unknown | -1.6512 | -3.1 | 0.00120209 |
| YIR038C | GTT1 | glutathione transferase | -1.6492 | -3.1 | 9.55407E-05 |
| YIR040C | | | -1.6405 | -3.1 | 0.0122417 |
| YNL173C | MDG1 | unknown | -1.6380 | -3.1 | 0.00548629 |
| YLR164W | | unknown | -1.6185 | -3.1 | 0.000293696 |
| YLR371W | ROM2 | signal transducer * | -1.6180 | -3.1 | 0.075157 |
| YOL083W | | unknown | -1.5882 | -3.0 | 0.000994279 |
| YLR130C | ZRT2 | low-affinity zinc ion transporter | -1.5797 | -3.0 | 0.0102896 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-------------|
| YIR036C | | unknown | -1.5775 | -3.0 | 0.0075153 |
| YOR273C | TPO4 | spermine transporter | -1.5768 | -3.0 | 0.0477474 |
| YGL145W | TIP20 | unknown | -1.5640 | -3.0 | 0.00343406 |
| YBR072W | HSP26 | unfolded protein binding | -1.5610 | -3.0 | 0.00139315 |
| YHR018C | ARG4 | argininosuccinate lyase | -1.5507 | -2.9 | 0.03339 |
| YBR067C | TIP1 | structural constituent of cell wall* | -1.5468 | -2.9 | 0.00537742 |
| YGR271W | SLH1 | RNA helicase | -1.5133 | -2.9 | 0.00143023 |
| YKL050C | | unknown | -1.5112 | -2.9 | 0.000526195 |
| YPL004C | LSP1 | protein kinase | -1.4955 | -2.8 | 0.00346151 |
| YGR146C | | unknown | -1.4940 | -2.8 | 0.00641131 |
| YDL232W | OST4 | protein binding | -1.4845 | -2.8 | 0.0198544 |
| YJR148W | BAT2 | branched-chain-amino-acid transaminase | -1.4750 | -2.8 | 0.0146444 |
| YJL039C | NUP192 | structural constituent of nuclear pore | -1.4678 | -2.8 | 0.0344339 |
| YBL013W | FMT1 | methionyl-tRNA formyltransferase | -1.4555 | -2.7 | 0.0120324 |
| YMR085W | | unknown | -1.4387 | -2.7 | 0.0656791 |
| YDR438W | | unknown | -1.4293 | -2.7 | 0.00396554 |
| YKR076W | ECM4 | unknown | -1.4290 | -2.7 | 0.00661339 |
| YAL043C | PTA1 | RNA binding | -1.4202 | -2.7 | 0.00367684 |
| YDR111C | ALT2 | transaminase | -1.4067 | -2.7 | 0.00859708 |
| YJL085W | EXO70 | protein binding | -1.4025 | -2.6 | 0.00894946 |
| YBR222C | PCS60 | AMP binding | -1.3972 | -2.6 | 0.00133507 |
| YOR290C | SNF2 | general RNA polymerase II transcription factor | -1.3777 | -2.6 | 0.061678 |
| YJR104C | SOD1 | copper, zinc superoxide dismutase | -1.3720 | -2.6 | 0.00125702 |
| YMR006C | PLB2 | lysophospholipase | -1.3660 | -2.6 | 0.0152239 |
| YMR041C | | unknown | -1.3607 | -2.6 | 0.0627898 |
| YLL053C | | unknown | -1.3583 | -2.6 | 0.12105 |
| YMR031C | | unknown | -1.3423 | -2.5 | 0.0204176 |
| YNL014W | HEF3 | ATPase * | -1.3380 | -2.5 | 0.000928258 |
| YPL278C | | unknown | -1.3358 | -2.5 | 0.000933247 |
| YFL006W | | | -1.3337 | -2.5 | 0.00970652 |
| YLR283W | | unknown | -1.3327 | -2.5 | 0.00560181 |
| YIL136W | OM45 | unknown | -1.3317 | -2.5 | 0.00925432 |
| YOR342C | | unknown | -1.3288 | -2.5 | 0.0217426 |
| YML008C | ERG6 | sterol 24-C-methyltransferase | -1.3130 | -2.5 | 0.00512992 |
| YGL205W | POX1 | acyl-CoA oxidase | -1.3047 | -2.5 | 0.00114115 |
| YNL208W | | unknown | -1.3030 | -2.5 | 0.0108054 |
| YDL215C | GDH2 | glutamate dehydrogenase | -1.2965 | -2.5 | 0.00240171 |
| YDR385W | EFT2 | translation elongation factor | -1.2947 | -2.5 | 0.00958742 |
| YBR094W | | unknown | -1.2908 | -2.4 | 0.0118839 |
| YER091C | MET6 | 5-methyltetrahydropteroyltriglutamate-homocysteine S-met | -1.2880 | -2.4 | 0.0012232 |
| YOR101W | RAS1 | GTPase | -1.2768 | -2.4 | 0.0246678 |
| YDR319C | | unknown | -1.2712 | -2.4 | 0.00046448 |
| YPL061W | ALD6 | 3-chloroallyl aldehyde dehydrogenase | -1.2535 | -2.4 | 0.0103137 |
| YDL085W | NDE2 | NADH dehydrogenase | -1.2443 | -2.4 | 0.00672484 |
| YGR247W | CPD1 | 2',3'-cyclic-nucleotide 3'-phosphodiesterase | -1.2383 | -2.4 | 0.00429145 |
| YGR008C | STF2 | unknown | -1.2238 | -2.3 | 0.00971904 |
| YAL067C | SEO1 | transporter | -1.2233 | -2.3 | 0.00757927 |
| YPL113C | | oxidoreductase | -1.2220 | -2.3 | 0.0205914 |
| YGR154C | GTO1 | unknown | -1.2203 | -2.3 | 0.0372931 |
| YGR056W | RSC1 | unknown | -1.2193 | -2.3 | 0.00918573 |
| YLR389C | STE23 | metallopeptidase | -1.2183 | -2.3 | 0.00519147 |
| YLR267W | BOP2 | unknown | -1.2130 | -2.3 | 0.00069634 |
| YNR001C | CIT1 | citrate (Si)-synthase | -1.2120 | -2.3 | 0.00125313 |
| YEL028W | | | -1.2077 | -2.3 | 0.0106174 |
| YGL062W | PYC1 | pyruvate carboxylase | -1.2017 | -2.3 | 0.00807525 |
| YGR161C | RTS3 | protein phosphatase type 2A | -1.1925 | -2.3 | 0.00124208 |
| YKR009C | FOX2 | 3-hydroxyacyl-CoA dehydrogenase | -1.1897 | -2.3 | 0.0332398 |
| YLR348C | DIC1 | dicarboxylic acid transporter | -1.1777 | -2.3 | 0.00787494 |
| YGL196W | | unknown | -1.1738 | -2.3 | 0.0163304 |
| YER124C | DSE1 | unknown | -1.1725 | -2.3 | 0.0582102 |
| YJL163C | | unknown | -1.1620 | -2.2 | 0.0160042 |
| YIL124W | AYR1 | acylglycerone-phosphate reductase | -1.1532 | -2.2 | 0.000707662 |
| YER121W | | | -1.1497 | -2.2 | 0.00201393 |
| YAR042W | SWH1 | phosphatidylinositol binding | -1.1483 | -2.2 | 0.0384143 |
| YMR214W | SCJ1 | chaperone binding | -1.1378 | -2.2 | 0.00542857 |
| YGR086C | PIL1 | protein kinase | -1.1333 | -2.2 | 0.00138338 |
| YAR040C | | | -1.1330 | -2.2 | 0.013239 |
| YMR072W | ABF2 | DNA binding | -1.1308 | -2.2 | 0.0142121 |
| YIL009W | FAA3 | long-chain-fatty-acid-CoA ligase | -1.1298 | -2.2 | 0.0118662 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YOL126C | MDH2 | L-malate dehydrogenase | -1.1280 | -2.2 | 0.00466172 |
| YMR303C | ADH2 | alcohol dehydrogenase | -1.1158 | -2.2 | 0.0242276 |
| YPL246C | RBD2 | unknown | -1.1157 | -2.2 | 0.0158778 |
| YHR096C | HXT5 | glucose transporter | -1.1155 | -2.2 | 0.031484 |
| YMR315W | | unknown | -1.1118 | -2.2 | 0.0450408 |
| YDR497C | ITR1 | myo-inositol transporter | -1.1070 | -2.2 | 0.00777437 |
| YOR133W | EFT1 | translation elongation factor | -1.1047 | -2.2 | 0.0332233 |
| YPR150W | | | -1.1033 | -2.1 | 0.0116281 |
| YOL058W | ARG1 | argininosuccinate synthase | -1.0945 | -2.1 | 0.0134545 |
| YGR052W | FMP48 | kinase | -1.0840 | -2.1 | 0.00176231 |
| YIL160C | POT1 | acetyl-CoA C-acyltransferase | -1.0828 | -2.1 | 0.0255099 |
| YMR246W | FAA4 | long-chain-fatty-acid-CoA ligase | -1.0808 | -2.1 | 0.00450923 |
| YEL060C | PRB1 | serine-type endopeptidase | -1.0758 | -2.1 | 0.0232746 |
| YJR073C | OPI3 | phosphatidyl-N-methylethanolamine N-methyltransferase | -1.0692 | -2.1 | 0.00285444 |
| YJL045W | | succinate dehydrogenase | -1.0680 | -2.1 | 0.0377457 |
| YOL110W | SHR5 | protein-cysteine S-palmitoleyltransferase | -1.0673 | -2.1 | 0.0170142 |
| YHR007C | ERG11 | sterol 14-demethylase | -1.0665 | -2.1 | 0.0151038 |
| YOR373W | NUD1 | structural constituent of cytoskeleton | -1.0662 | -2.1 | 0.0256083 |
| YBR046C | ZTA1 | unknown | -1.0640 | -2.1 | 0.00188264 |
| YER084W | | | -1.0623 | -2.1 | 0.000970999 |
| YNL057W | | | -1.0615 | -2.1 | 0.0259222 |
| YLR355C | ILV5 | ketol-acid reductoisomerase | -1.0612 | -2.1 | 0.00223063 |
| YDL046W | NPC2 | unknown | -1.0610 | -2.1 | 0.00236527 |
| YMR191W | SPG5 | unknown | -1.0607 | -2.1 | 0.00187758 |
| YGR175C | ERG1 | squalene monooxygenase * | -1.0603 | -2.1 | 0.0279659 |
| YPL028W | ERG10 | acetyl-CoA C-acetyltransferase | -1.0572 | -2.1 | 0.0327046 |
| YJL088W | ARG3 | ornithine carbamoyltransferase | -1.0455 | -2.1 | 0.0143854 |
| YMR152W | YIM1 | unknown | -1.0428 | -2.1 | 0.0119463 |
| YOL053C-A | | | -1.0337 | -2.0 | 0.0101263 |
| YJL067W | | | -1.0325 | -2.0 | 0.0457892 |
| YNL115C | | unknown | -1.0318 | -2.0 | 0.00478682 |
| YNL195C | | unknown | -1.0278 | -2.0 | 0.0280353 |
| YPL147W | PXA1 | ATPase , coupled to transmembrane movement of substance | -1.0272 | -2.0 | 0.0185403 |
| YEL071W | DLD3 | D-lactate dehydrogenase | -1.0162 | -2.0 | 0.0501081 |
| YKL217W | JEN1 | lactate transporter | -1.0117 | -2.0 | 0.0502041 |
| YMR160W | | unknown | -1.0105 | -2.0 | 0.0206442 |
| YBL016W | FUS3 | MAP kinase | -1.0055 | -2.0 | 0.0239609 |
| YIL047C | SYG1 | unknown | -1.0007 | -2.0 | 0.0106308 |
| YLR369W | SSQ1 | unfolded protein binding | -0.9982 | -2.0 | 0.164697 |
| YDR039C | ENA2 | ATPase , coupled to transmembrane movement of ions, pho | -0.9973 | -2.0 | 0.0332456 |
| YNL009W | IDP3 | isocitrate dehydrogenase | -0.9895 | -2.0 | 0.0191983 |
| YMR083W | ADH3 | alcohol dehydrogenase | -0.9872 | -2.0 | 0.0232675 |
| YDR046C | BAP3 | amino acid transporter | -0.9850 | -2.0 | 0.0161152 |
| YCR004C | YCP4 | electron carrier | -0.9833 | -2.0 | 0.00378332 |
| YHR001W | OSH7 | oxysterol binding | -0.9828 | -2.0 | 0.0115582 |
| YMR251W-A | HOR7 | unknown | -0.9800 | -2.0 | 0.00778365 |
| YGL184C | STR3 | cystathionine beta-lyase | -0.9758 | -2.0 | 0.000968937 |
| YMR297W | PRC1 | carboxypeptidase C | -0.9695 | -2.0 | 0.00983392 |
| YIL177C | | helicase | -0.9693 | -2.0 | 0.00539022 |
| YLR081W | GAL2 | glucose transporter | -0.9660 | -2.0 | 0.0561172 |
| YOR229W | WTM2 | transcription corepressor | -0.9615 | -1.9 | 0.0126279 |
| YLR320W | MMS22 | unknown | -0.9605 | -1.9 | 0.0182108 |
| YPR139C | VPS66 | unknown | -0.9583 | -1.9 | 0.00745739 |
| YMR056C | AAC1 | ATP:ADP antiporter | -0.9543 | -1.9 | 0.0320628 |
| YJL026W | RNR2 | ribonucleoside-diphosphate reductase | -0.9527 | -1.9 | 0.00274444 |
| YHR208W | BAT1 | branched-chain-amino-acid transaminase | -0.9523 | -1.9 | 0.0278678 |
| YDR033W | MRH1 | unknown | -0.9478 | -1.9 | 0.0036846 |
| YGL156W | AMS1 | alpha-mannosidase | -0.9452 | -1.9 | 0.0352944 |
| YDL073W | | unknown | -0.9368 | -1.9 | 0.0480045 |
| YHR092C | HXT4 | glucose transporter | -0.9333 | -1.9 | 0.0327426 |
| YJR008W | | unknown | -0.9323 | -1.9 | 0.00354344 |
| YBL075C | SSA3 | ATPase * | -0.9300 | -1.9 | 0.0127402 |
| YHR215W | PHO12 | acid phosphatase | -0.9293 | -1.9 | 0.00354773 |
| YIL115C | NUP159 | structural molecule | -0.9272 | -1.9 | 0.00832438 |
| YOL086C | ADH1 | alcohol dehydrogenase | -0.9160 | -1.9 | 0.0153947 |
| YLR467W | YRF1-5 | DNA helicase | -0.9117 | -1.9 | 0.0144548 |
| YOR375C | GDH1 | glutamate dehydrogenase | -0.9117 | -1.9 | 0.0909276 |
| YOR348C | PUT4 | L-proline permease | -0.9108 | -1.9 | 0.0414861 |
| YJR107W | | lipase | -0.9088 | -1.9 | 0.121761 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|---|----------------------------|-------------|-------------|
| YDL182W | LYS20 | homocitrate synthase | -0.9077 | -1.9 | 0.000343853 |
| YIL166C | | transporter | -0.8972 | -1.9 | 0.0112094 |
| YDL173W | | unknown | -0.8942 | -1.9 | 0.00417624 |
| YCR047C | BUD23 | S-adenosylmethionine-dependent methyltransferase | -0.8927 | -1.9 | 0.0126782 |
| YPL049C | DIG1 | DNA binding | -0.8923 | -1.9 | 0.0125409 |
| YGR180C | RNR4 | ribonucleoside-diphosphate reductase | -0.8910 | -1.9 | 0.0207537 |
| YLR327C | TMA10 | unknown | -0.8853 | -1.8 | 0.00970216 |
| YCL004W(1) | | | -0.8823 | -1.8 | 0.0139477 |
| YIR002C | MPH1 | RNA helicase | -0.8813 | -1.8 | 0.00902535 |
| YLL052C | AQY2 | water channel | -0.8812 | -1.8 | 0.0206034 |
| YLR174W | IDP2 | isocitrate dehydrogenase | -0.8795 | -1.8 | 0.019632 |
| YMR195W | ICY1 | unknown | -0.8792 | -1.8 | 0.00503988 |
| YFR036W-A | | | -0.8792 | -1.8 | 0.0451375 |
| YDR032C | PST2 | unknown | -0.8775 | -1.8 | 0.00213569 |
| YBR144C | | | -0.8765 | -1.8 | 0.0428772 |
| YOR095C | RKI1 | ribose-5-phosphate isomerase | -0.8748 | -1.8 | 0.00791865 |
| YPR010C | RPA135 | DNA-directed RNA polymerase | -0.8745 | -1.8 | 0.0288493 |
| YMR116C | ASC1 | unknown | -0.8728 | -1.8 | 0.0020079 |
| YIL119C | RPI1 | small GTPase regulator | -0.8722 | -1.8 | 0.0277949 |
| YIL033C | BCY1 | cAMP-dependent protein kinase | -0.8713 | -1.8 | 0.0576105 |
| YKL199C | | | -0.8675 | -1.8 | 0.0202703 |
| YKL033W | | unknown | -0.8670 | -1.8 | 0.0244836 |
| YJL219W | HXT9 | glucose transporter | -0.8655 | -1.8 | 0.0332624 |
| YJL130C | URA2 | carbamoyl-phosphate synthase (glutamine-hydrolyzing) * | -0.8623 | -1.8 | 0.0180221 |
| YFR044C | | unknown | -0.8602 | -1.8 | 0.0224477 |
| YHR094C | HXT1 | glucose transporter | -0.8542 | -1.8 | 0.0336488 |
| YMR139W | RIM11 | protein serine/threonine kinase | -0.8488 | -1.8 | 0.29394 |
| YOR054C | VHS3 | phosphopantothienoylcysteine decarboxylase | -0.8488 | -1.8 | 0.0502112 |
| YBR068C | BAP2 | amino acid transporter | -0.8447 | -1.8 | 0.00322697 |
| YLR414C | | unknown | -0.8437 | -1.8 | 0.00244411 |
| YOL153C | | unknown | -0.8388 | -1.8 | 0.0439778 |
| YER084W-A | | | -0.8360 | -1.8 | 0.221601 |
| YFL011W | HXT10 | glucose transporter | -0.8352 | -1.8 | 0.002615 |
| YAL056W | GPB2 | signal transducer | -0.8345 | -1.8 | 0.0003491 |
| YIR039C | YPS6 | aspartic-type endopeptidase | -0.8338 | -1.8 | 0.0522286 |
| YMR250W | GAD1 | glutamate decarboxylase | -0.8335 | -1.8 | 0.0295853 |
| YNL008C | ASI3 | ubiquitin-protein ligase | -0.8330 | -1.8 | 0.0218619 |
| YIR037W | HYR1 | thioredoxin peroxidase | -0.8285 | -1.8 | 0.00356745 |
| YIL170W | HXT12 | unknown | -0.8280 | -1.8 | 0.0527226 |
| YOR031W | CRS5 | copper ion binding | -0.8278 | -1.8 | 0.00849453 |
| YNL239W | LAP3 | transcription regulator | -0.8267 | -1.8 | 0.104249 |
| YMR164C | MSS11 | specific RNA polymerase II transcription factor | -0.8260 | -1.8 | 0.0892817 |
| YML118W | NGL3 | endonuclease | -0.8248 | -1.8 | 0.00521392 |
| YPL103C | FMP30 | unknown | -0.8238 | -1.8 | 0.0799272 |
| YKL057C | NUP120 | structural molecule | -0.8182 | -1.8 | 0.172077 |
| YBR025C | | unknown | -0.8165 | -1.8 | 0.00703013 |
| YGL017W | ATE1 | arginyltransferase | -0.8155 | -1.8 | 0.0192379 |
| YBR054W | YRO2 | unknown | -0.8132 | -1.8 | 0.0395992 |
| YGR130C | | unknown | -0.8085 | -1.8 | 0.0177596 |
| YNL134C | | alcohol dehydrogenase | -0.8077 | -1.8 | 0.0099102 |
| YBR083W | TEC1 | specific RNA polymerase II transcription factor | -0.8068 | -1.7 | 0.0207251 |
| YPL062W | | | -0.8052 | -1.7 | 0.10305 |
| YJL225C | | helicase | -0.8017 | -1.7 | 0.0174451 |
| YKL005C | BYE1 | transcriptional elongation regulator | -0.8017 | -1.7 | 0.0042893 |
| YER138C | | | -0.8013 | -1.7 | 0.113771 |
| YLL066W-A | | | -0.7983 | -1.7 | 0.0394687 |
| YPL254W | HF11 | transcription coactivator * | -0.7972 | -1.7 | 0.189492 |
| YHL034C | SBP1 | RNA binding | -0.7935 | -1.7 | 0.000827879 |
| YML010W-B | | | -0.7923 | -1.7 | 0.124338 |
| YPL264C | | unknown | -0.7920 | -1.7 | 0.000450074 |
| YLR306W | UBC12 | NEDD8 conjugating enzyme | -0.7910 | -1.7 | 0.000975033 |
| YNR050C | LYS9 | saccharopine dehydrogenase | -0.7850 | -1.7 | 0.00158771 |
| YJR139C | HOM6 | homoserine dehydrogenase | -0.7848 | -1.7 | 0.0133602 |
| YPR012W | | | -0.7787 | -1.7 | 0.0489419 |
| YLR356W | | unknown | -0.7785 | -1.7 | 0.0226944 |
| YLL020C | | | -0.7768 | -1.7 | 0.00430602 |
| YOL123W | HRP1 | RNA binding | -0.7747 | -1.7 | 0.0259376 |
| YLL016W | SDC25 | Ras guanyl-nucleotide exchange factor | -0.7745 | -1.7 | 0.00764971 |
| YDR038C | ENA5 | ATPase , coupled to transmembrane movement of ions, pho | -0.7725 | -1.7 | 0.0714362 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YPL109C | | unknown | -0.7715 | -1.7 | 0.025346 |
| YNL200C | | unknown | -0.7687 | -1.7 | 0.0135264 |
| YAR018C | KIN3 | protein kinase | -0.7663 | -1.7 | 0.0126639 |
| YMR318C | ADH6 | alcohol dehydrogenase | -0.7657 | -1.7 | 0.0238493 |
| YMR206W | | unknown | -0.7625 | -1.7 | 0.00165901 |
| YDR300C | PRO1 | glutamate 5-kinase | -0.7597 | -1.7 | 0.0259768 |
| YOL065C | INP54 | inositol-polyphosphate 5-phosphatase | -0.7592 | -1.7 | 0.0063353 |
| YMR251W | GTO3 | unknown | -0.7585 | -1.7 | 0.14233 |
| YKL197C | PEX1 | ATPase | -0.7573 | -1.7 | 0.146408 |
| YOR184W | SER1 | phosphoserine transaminase | -0.7538 | -1.7 | 0.250373 |
| YBR286W | APE3 | aminopeptidase | -0.7537 | -1.7 | 0.0292206 |
| YBL113W-A | | | -0.7518 | -1.7 | 0.0448722 |
| YNL268W | LYP1 | basic amino acid transporter | -0.7517 | -1.7 | 0.0364305 |
| YOR336W | KRE5 | UDP-glucose:glycoprotein glucosyltransferase | -0.7507 | -1.7 | 0.0484315 |
| YBR244W | GPX2 | glutathione peroxidase | -0.7505 | -1.7 | 0.00137412 |
| YIR015W | RPR2 | ribonuclease P | -0.7500 | -1.7 | 0.0316138 |
| YBL042C | FUI1 | uridine transporter | -0.7498 | -1.7 | 0.000486736 |
| YDL169C | UGX2 | unknown | -0.7495 | -1.7 | 0.00759826 |
| YCR056W | | | -0.7452 | -1.7 | 0.0252701 |
| YOL147C | PEX11 | unknown | -0.7450 | -1.7 | 0.041068 |
| YLR214W | FRE1 | ferric-chelate reductase | -0.7440 | -1.7 | 0.00501358 |
| YPR173C | VPS4 | ATPase | -0.7410 | -1.7 | 0.0300097 |
| YBL014C | RRN6 | RNA polymerase I transcription factor | -0.7378 | -1.7 | 0.0319721 |
| YPL119C | DBP1 | RNA helicase | -0.7348 | -1.7 | 0.263192 |
| YMR110C | HFD1 | 3-chloroallyl aldehyde dehydrogenase | -0.7338 | -1.7 | 0.00668068 |
| YJL042W | MHP1 | structural constituent of cytoskeleton | -0.7317 | -1.7 | 0.00112377 |
| YGR155W | CYS4 | cystathionine beta-synthase | -0.7302 | -1.7 | 0.00020141 |
| YDL171C | GLT1 | glutamate synthase (NADH) | -0.7295 | -1.7 | 0.0297633 |
| YPL018W | CTF19 | protein binding | -0.7293 | -1.7 | 0.298707 |
| YNL246W | VPS75 | unknown | -0.7287 | -1.7 | 0.00399544 |
| YOL018C | TLG2 | v-SNARE * | -0.7280 | -1.7 | 0.236142 |
| YBR139W | | serine hydrolase * | -0.7277 | -1.7 | 0.0186737 |
| YBR020W | GAL1 | galactokinase | -0.7275 | -1.7 | 0.030461 |
| YKL157W | APE2 | leucyl aminopeptidase | -0.7245 | -1.7 | 0.0142352 |
| YEL063C | CAN1 | basic amino acid transporter | -0.7240 | -1.7 | 0.0502109 |
| YNL015W | PBI2 | endopeptidase inhibitor | -0.7208 | -1.6 | 0.0767774 |
| YMR019W | STB4 | DNA binding | -0.7207 | -1.6 | 0.0519094 |
| YMR114C | | unknown | -0.7203 | -1.6 | 0.0150624 |
| YCR011C | ADP1 | ATPase , coupled to transmembrane movement of substance | -0.7202 | -1.6 | 0.0508017 |
| YHR074W | QNS1 | hydrolase , acting on carbon-nitrogen (but not peptide) bonds | -0.7202 | -1.6 | 0.120026 |
| YCL049C | | unknown | -0.7200 | -1.6 | 0.00646251 |
| YEL058W | PCM1 | phosphoacetylglucosamine mutase | -0.7187 | -1.6 | 0.00194544 |
| YCRX02C | | | -0.7187 | -1.6 | 0.0384456 |
| YPR025C | CCL1 | general RNA polymerase II transcription factor | -0.7162 | -1.6 | 0.0412672 |
| YLR450W | HMG2 | hydroxymethylglutaryl-CoA reductase (NADPH) | -0.7160 | -1.6 | 0.154623 |
| YGR141W | VPS62 | unknown | -0.7150 | -1.6 | 0.0198467 |
| YLL002W | RTT109 | unknown | -0.7147 | -1.6 | 0.00725145 |
| YBR265W | TSC10 | oxidoreductase | -0.7138 | -1.6 | 0.00954183 |
| YPL186C | UIP4 | unknown | -0.7137 | -1.6 | 0.0223697 |
| Q0110 | BI2 | nuclease | -0.7120 | -1.6 | 0.0261979 |
| YDR015C | | | -0.7098 | -1.6 | 0.0546078 |
| YMR102C | | unknown | -0.7095 | -1.6 | 0.0849514 |
| YNL270C | ALP1 | basic amino acid transporter | -0.7075 | -1.6 | 0.0330172 |
| YBR053C | | unknown | -0.7068 | -1.6 | 0.00179155 |
| YEL011W | GLC3 | 1,4-alpha-glucan branching enzyme | -0.7062 | -1.6 | 0.00120619 |
| YNL307C | MCK1 | glycogen synthase kinase | -0.7058 | -1.6 | 0.034878 |
| YLR460C | | unknown | -0.7040 | -1.6 | 0.0528768 |
| YER130C | | unknown | -0.7030 | -1.6 | 0.0164337 |
| YDL078C | MDH3 | L-malate dehydrogenase | -0.7028 | -1.6 | 0.0198745 |
| YDL222C | FMP45 | unknown | -0.7027 | -1.6 | 0.00370794 |
| YER172C | BRR2 | RNA splicing factor | -0.7022 | -1.6 | 0.0715497 |
| YJL116C | NCA3 | unknown | -0.6948 | -1.6 | 0.0499002 |
| YNL168C | FMP41 | unknown | -0.6930 | -1.6 | 0.0106248 |
| YPR013C | | unknown | -0.6927 | -1.6 | 0.00832789 |
| YMR108W | ILV2 | acetolactate synthase * | -0.6835 | -1.6 | 0.0118143 |
| YAL055W | PEX22 | unknown | -0.6795 | -1.6 | 0.0141397 |
| YIL071C | PCI8 | unknown | -0.6792 | -1.6 | 0.00119162 |
| YMR119W-A | | | -0.6772 | -1.6 | 0.00780916 |
| YCR093W | CDC39 | 3'-5'-exoribonuclease | -0.6765 | -1.6 | 0.00752911 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YJR112W | NNF1 | unknown | -0.6753 | -1.6 | 0.00507128 |
| YLR143W | | unknown | -0.6752 | -1.6 | 0.0924621 |
| YNL280C | ERG24 | delta14-sterol reductase | -0.6750 | -1.6 | 0.133412 |
| YOR165W | SEY1 | unknown | -0.6742 | -1.6 | 0.112196 |
| YCRX15W | | | -0.6735 | -1.6 | 0.0498245 |
| YKL123W | | | -0.6733 | -1.6 | 0.0326957 |
| YBR131W | CCZ1 | guanyl-nucleotide exchange factor | -0.6727 | -1.6 | 0.0176449 |
| YFR024C-A | LSB3 | unknown | -0.6717 | -1.6 | 0.0190464 |
| YBR080C | SEC18 | ATPase | -0.6697 | -1.6 | 0.00238185 |
| YBL067C | UBP13 | ubiquitin-specific protease | -0.6693 | -1.6 | 0.008725 |
| YNL067W | RPL9B | structural constituent of ribosome | -0.6690 | -1.6 | 0.0126196 |
| YDR129C | SAC6 | protein binding | -0.6660 | -1.6 | 0.0821353 |
| YPL037C | EGD1 | unfolded protein binding | -0.6647 | -1.6 | 0.0293449 |
| YHR097C | | unknown | -0.6608 | -1.6 | 0.0185694 |
| YLR438W | CAR2 | ornithine-oxo-acid transaminase | -0.6593 | -1.6 | 0.0153655 |
| YGR067C | | unknown | -0.6565 | -1.6 | 0.023154 |
| YER178W | PDA1 | pyruvate dehydrogenase | -0.6545 | -1.6 | 0.0220401 |
| YHR072W | ERG7 | lanosterol synthase | -0.6535 | -1.6 | 0.0129567 |
| YOL103W | ITR2 | myo-inositol transporter | -0.6495 | -1.6 | 0.121942 |
| YKR084C | HBS1 | unknown | -0.6488 | -1.6 | 0.000758252 |
| YPR080W | TEF1 | translation elongation factor | -0.6478 | -1.6 | 0.166537 |
| YOR040W | GLO4 | hydroxyacylglutathione hydrolase | -0.6477 | -1.6 | 0.111901 |
| YPL134C | ODC1 | intracellular transporter | -0.6463 | -1.6 | 0.00333852 |
| YML072C | TCB3 | lipid binding | -0.6462 | -1.6 | 0.0119512 |
| YPL074W | YTA6 | ATPase | -0.6455 | -1.6 | 0.0791675 |
| YJL172W | CPS1 | Gly-X carboxypeptidase | -0.6432 | -1.6 | 0.0579268 |
| YGL059W | | protein kinase | -0.6397 | -1.6 | 0.0254145 |
| YCR062W | | | -0.6383 | -1.6 | 0.000857556 |
| YEL016C | NPP2 | nucleoside-triphosphatase * | -0.6383 | -1.6 | 0.110499 |
| YGL055W | OLE1 | stearyl-CoA 9-desaturase | -0.6365 | -1.6 | 0.0514347 |
| YDR247W | VHS1 | protein kinase | -0.6360 | -1.6 | 0.102614 |
| YGL161C | YIP5 | Rab GTPase binding | -0.6357 | -1.6 | 0.0342868 |
| YDR160W | SSY1 | amino acid binding | -0.6303 | -1.5 | 0.0671155 |
| YER020W | GPA2 | GTPase | -0.6288 | -1.5 | 0.00348378 |
| YLR237W | THI7 | thiamin transporter | -0.6282 | -1.5 | 0.0478375 |
| YCL013W | | | -0.6278 | -1.5 | 0.00123192 |
| YDR506C | | unknown | -0.6273 | -1.5 | 0.132163 |
| YDR246W | TRS23 | unknown | -0.6272 | -1.5 | 0.0978044 |
| YDL179W | PCL9 | cyclin-dependent protein kinase | -0.6263 | -1.5 | 0.0274064 |
| YNL321W | | unknown | -0.6262 | -1.5 | 0.107064 |
| YNR016C | ACC1 | acetyl-CoA carboxylase * | -0.6242 | -1.5 | 0.0356062 |
| YJR130C | STR2 | cystathionine gamma-synthase | -0.6238 | -1.5 | 0.00401844 |
| YNL093W | YPT53 | GTPase | -0.6232 | -1.5 | 0.10927 |
| YLR178C | TFS1 | lipid binding | -0.6230 | -1.5 | 0.00316436 |
| YER103W | SSA4 | ATPase * | -0.6228 | -1.5 | 0.0342273 |
| YIL094C | LYS12 | homoisocitrate dehydrogenase | -0.6215 | -1.5 | 0.0524943 |
| YFR047C | BNA6 | nicotinate-nucleotide diphosphorylase (carboxylating) | -0.6182 | -1.5 | 0.0108959 |
| YGR134W | CAF130 | unknown | -0.6155 | -1.5 | 0.0326339 |
| YOL165C | AAD15 | aryl-alcohol dehydrogenase | -0.6135 | -1.5 | 0.0442084 |
| YDL200C | MGT1 | methylated-DNA-[protein]-cysteine S-methyltransferase | -0.6120 | -1.5 | 0.0588737 |
| YMR238W | DFG5 | mannosidase | -0.6108 | -1.5 | 0.0255659 |
| YCR086W | CSM1 | unknown | -0.6107 | -1.5 | 0.0445425 |
| YJL147C | | unknown | -0.6090 | -1.5 | 0.00069081 |
| YFR041C | ERJ5 | unknown | -0.6088 | -1.5 | 0.00303969 |
| YLR093C | NYV1 | v-SNARE | -0.6087 | -1.5 | 0.00229792 |
| YDL239C | ADY3 | protein binding | -0.6072 | -1.5 | 0.0512154 |
| YPR003C | | unknown | -0.6070 | -1.5 | 0.0254132 |
| YBR097W | VPS15 | protein serine/threonine kinase | -0.6068 | -1.5 | 0.171751 |
| YLR205C | HMX1 | heme binding | -0.6057 | -1.5 | 0.00028184 |
| YML010W-A | | | -0.6057 | -1.5 | 0.11892 |
| YPR133C | SPN1 | RNA polymerase II transcription elongation factor | -0.6028 | -1.5 | 0.00158878 |
| YMR106C | YKU80 | RNA binding | -0.6020 | -1.5 | 0.0711883 |
| YKL117W | SBA1 | chaperone binding | -0.6017 | -1.5 | 0.0136933 |
| YIL158W | | unknown | -0.5998 | -1.5 | 0.00198558 |
| YHL010C | | nuclear localization sequence binding | -0.5982 | -1.5 | 0.0633709 |
| YOR019W | | unknown | -0.5952 | -1.5 | 0.0734784 |
| YJR039W | | unknown | -0.5950 | -1.5 | 0.0469764 |
| YKL210W | UBA1 | ubiquitin activating enzyme | -0.5945 | -1.5 | 0.00132668 |
| YDR179W-A | | unknown | -0.5945 | -1.5 | 0.0388303 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-------------|
| YPR108W | RPN7 | structural molecule | -0.5937 | -1.5 | 0.0188019 |
| YCR102C | | unknown | -0.5933 | -1.5 | 0.00549815 |
| YPR015C | | unknown | -0.5923 | -1.5 | 0.0167209 |
| YOL062C | APM4 | unknown | -0.5915 | -1.5 | 0.149685 |
| YOR304W | ISW2 | ATPase | -0.5913 | -1.5 | 0.0275905 |
| YDR433W | | | -0.5913 | -1.5 | 0.0854852 |
| YJR099W | YUH1 | ubiquitin-specific protease | -0.5910 | -1.5 | 0.00976385 |
| YIL046W | MET30 | protein binding | -0.5898 | -1.5 | 0.112747 |
| YPR082C | DIB1 | RNA splicing factor | -0.5887 | -1.5 | 0.0189485 |
| YDR249C | | unknown | -0.5885 | -1.5 | 0.0414751 |
| YDR358W | GGA1 | ubiquitin binding | -0.5880 | -1.5 | 0.00056417 |
| YDL185W | TFP1 | hydrogen-transporting ATPase , rotational mechanism* | -0.5880 | -1.5 | 0.0597065 |
| YAR071W | PHO11 | acid phosphatase | -0.5878 | -1.5 | 0.00744237 |
| YGR256W | GND2 | phosphogluconate dehydrogenase | -0.5865 | -1.5 | 0.0455373 |
| YPL154C | PEP4 | endopeptidase * | -0.5850 | -1.5 | 0.0292443 |
| YOR192C | | transporter | -0.5837 | -1.5 | 0.00113163 |
| YJR066W | TOR1 | protein binding | -0.5820 | -1.5 | 0.0123851 |
| YJR131W | MNS1 | mannosyl-oligosaccharide 1,2-alpha-mannosidase | -0.5807 | -1.5 | 0.00633896 |
| YLR058C | SHM2 | glycine hydroxymethyltransferase | -0.5807 | -1.5 | 0.0883191 |
| YHR008C | SOD2 | manganese superoxide dismutase | -0.5795 | -1.5 | 0.00554771 |
| YGL164C | YRB30 | protein binding | -0.5788 | -1.5 | 0.00457977 |
| YDL032W | | | -0.5767 | -1.5 | 0.0595341 |
| YLR088W | GAA1 | GPI-anchor transamidase | -0.5762 | -1.5 | 0.0886485 |
| YMR258C | | unknown | -0.5760 | -1.5 | 0.000731584 |
| YDR285W | ZIP1 | chromatin binding | -0.5750 | -1.5 | 0.0192479 |
| YGR121C | MEP1 | ammonium transporter | -0.5723 | -1.5 | 0.0646348 |
| YJL132W | | unknown | -0.5707 | -1.5 | 0.0178218 |
| YOR175C | | O-acyltransferase | -0.5707 | -1.5 | 0.0125068 |
| YGL245W | GUS1 | glutamate-tRNA ligase | -0.5705 | -1.5 | 0.0476166 |
| YPR053C | | | -0.5700 | -1.5 | 0.036026 |
| YPL230W | | unknown | -0.5697 | -1.5 | 0.165868 |
| YLR401C | DUS3 | tRNA dihydrouridine synthase | -0.5692 | -1.5 | 0.114528 |
| YDR406W | PDR15 | ATPase , coupled to transmembrane movement of substance | -0.5678 | -1.5 | 0.161776 |
| YNR059W | MNT4 | alpha-1,3-mannosyltransferase | -0.5677 | -1.5 | 0.182102 |
| YML133C | | helicase | -0.5668 | -1.5 | 0.128108 |
| YOR356W | | oxidoreductase | -0.5665 | -1.5 | 0.249975 |
| YAR014C | BUD14 | protein phosphatase type 1 regulator | -0.5660 | -1.5 | 0.0107293 |
| YLR120C | YPS1 | aspartic-type endopeptidase | -0.5657 | -1.5 | 0.0716904 |
| YPL232W | SSO1 | t-SNARE | -0.5645 | -1.5 | 0.250982 |
| YDL199C | | unknown | -0.5638 | -1.5 | 0.0701341 |
| YBR214W | SDS24 | unknown | -0.5633 | -1.5 | 0.0245915 |
| YOL082W | ATG19 | protein binding | -0.5623 | -1.5 | 0.000890104 |
| YIL093C | RSM25 | structural constituent of ribosome | -0.5602 | -1.5 | 0.156624 |
| YLR365W | | | -0.5560 | -1.5 | 0.0871798 |
| YJL070C | | unknown | -0.5557 | -1.5 | 0.00281013 |
| YPL256C | CLN2 | cyclin-dependent protein kinase | -0.5548 | -1.5 | 0.183438 |
| YJL186W | MNN5 | alpha-1,2-mannosyltransferase | -0.5547 | -1.5 | 0.0550634 |
| YCR045C | | serine-type peptidase | -0.5542 | -1.5 | 0.0264241 |
| Q0105 | COB | ubiquinol-cytochrome-c reductase | -0.5542 | -1.5 | 0.0669248 |
| YAR043C | | | -0.5518 | -1.5 | 0.040718 |
| YOR259C | RPT4 | ATPase * | -0.5517 | -1.5 | 0.0198706 |
| YKL020C | SPT23 | DNA binding | -0.5515 | -1.5 | 0.142356 |
| YLR173W | | unknown | -0.5507 | -1.5 | 0.071563 |
| YLR219W | MSC3 | unknown | -0.5490 | -1.5 | 0.077245 |
| YLR337W | | | -0.5488 | -1.5 | 0.121434 |
| YLR084C | RAX2 | unknown | -0.5488 | -1.5 | 0.0751995 |
| YMR038C | CCS1 | superoxide dismutase copper chaperone | -0.5482 | -1.5 | 0.0136339 |
| YNL135C | FPR1 | peptidyl-prolyl cis-trans isomerase | -0.5472 | -1.5 | 0.00331367 |
| YJL164C | TPK1 | protein serine/threonine kinase | -0.5472 | -1.5 | 0.0462802 |
| YBL108W | | | -0.5465 | -1.5 | 0.0387518 |
| YDL144C | | unknown | -0.5463 | -1.5 | 0.0163463 |
| YIL082W | | | -0.5457 | -1.5 | 0.00657654 |
| YJR009C | TDH2 | glyceraldehyde-3-phosphate dehydrogenase | -0.5440 | -1.5 | 0.447956 |
| YJL024C | APS3 | unknown | -0.5402 | -1.5 | 0.00762379 |
| YKL144C | RPC25 | DNA-directed RNA polymerase | -0.5398 | -1.5 | 0.224644 |
| YOR215C | | unknown | -0.5393 | -1.5 | 0.0124647 |
| YBR243C | ALG7 | UDP-N-acetylglucosamine-dolichyl-phosphate N-acetylglucosaminyl transferase | -0.5392 | -1.5 | 0.0021535 |
| YHR136C | SPL2 | enzyme inhibitor | -0.5380 | -1.5 | 0.0230976 |
| YML065W | ORC1 | ATPase * | -0.5367 | -1.5 | 0.0471695 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-------------|
| YDL218W | | unknown | -0.5367 | -1.5 | 0.0116372 |
| YIL055C | | unknown | -0.5363 | -1.5 | 0.0862143 |
| YBR118W | TEF2 | translation elongation factor | -0.5360 | -1.4 | 0.0939697 |
| YPR032W | SRO7 | unknown | -0.5360 | -1.4 | 0.0133945 |
| YKR067W | GPT2 | glycerol-3-phosphate O-acyltransferase | -0.5353 | -1.4 | 0.012607 |
| YLL032C | | unknown | -0.5327 | -1.4 | 0.00509479 |
| YGL153W | PEX14 | protein binding | -0.5327 | -1.4 | 0.107384 |
| YDL086W | | carboxymethylenebutenolidase | -0.5312 | -1.4 | 0.0251987 |
| YBR069C | TAT1 | amino acid transporter | -0.5292 | -1.4 | 8.83079E-05 |
| YBR280C | | unknown | -0.5273 | -1.4 | 0.00187071 |
| YER023W | PRO3 | pyrroline-5-carboxylate reductase | -0.5263 | -1.4 | 0.025616 |
| YBR092C | PHO3 | acid phosphatase | -0.5255 | -1.4 | 0.0417088 |
| YNL170W | | | -0.5253 | -1.4 | 0.175941 |
| YKR049C | FMP46 | oxidoreductase | -0.5240 | -1.4 | 0.0619188 |
| YCRX18C | | | -0.5223 | -1.4 | 0.0791725 |
| YMR262W | | unknown | -0.5222 | -1.4 | 0.00984903 |
| Q0075 | AI5 BETA | unknown | -0.5197 | -1.4 | 0.0989434 |
| YPL193W | RSA1 | unknown | -0.5192 | -1.4 | 0.139014 |
| YBR093C | PHO5 | acid phosphatase * | -0.5188 | -1.4 | 0.0818423 |
| YMR052C-A | | | -0.5187 | -1.4 | 0.0704699 |
| YAL036C | RBG1 | GTP binding | -0.5187 | -1.4 | 0.0896977 |
| YDR417C | | | -0.5183 | -1.4 | 0.252248 |
| YNL212W | VID27 | unknown | -0.5167 | -1.4 | 0.0611748 |
| YIL083C | | phosphopantothenate--cysteine ligase | -0.5155 | -1.4 | 0.0512875 |
| YLR117C | CLF1 | protein binding | -0.5148 | -1.4 | 0.00842562 |
| YOR285W | | unknown | -0.5145 | -1.4 | 0.0897088 |
| YLR070C | XYL2 | D-xylulose reductase | -0.5133 | -1.4 | 0.0887707 |
| YPR104C | FHL1 | transcription factor | -0.5132 | -1.4 | 0.20462 |
| YJR106W | ECM27 | unknown | -0.5127 | -1.4 | 0.0295507 |
| YJL143W | TIM17 | protein transporter | -0.5127 | -1.4 | 0.220021 |
| YJL076W | NET1 | rDNA binding | -0.5123 | -1.4 | 0.000427566 |
| YAL044C | GCV3 | glycine dehydrogenase | -0.5103 | -1.4 | 0.143108 |
| YJL210W | PEX2 | protein binding | -0.5100 | -1.4 | 0.00431764 |
| YGL002W | ERP6 | unknown | -0.5078 | -1.4 | 0.00854339 |
| YIL112W | HOS4 | NAD-dependent histone deacetylase * | -0.5077 | -1.4 | 0.013199 |
| YNL163C | RIA1 | GTPase | -0.5077 | -1.4 | 0.0706703 |
| YMR129W | POM152 | structural molecule | -0.5065 | -1.4 | 0.119404 |
| YMR301C | ATM1 | ATPase , coupled to transmembrane movement of substanc | -0.5062 | -1.4 | 0.222795 |
| YCR088W | ABP1 | protein binding | -0.5060 | -1.4 | 0.033444 |
| YDL126C | CDC48 | ATPase | -0.5055 | -1.4 | 0.0979334 |
| YHL034W-A | | | -0.5050 | -1.4 | 0.0825645 |
| YMR054W | STV1 | hydrogen-transporting ATPase , rotational mechanism | -0.5048 | -1.4 | 0.0371778 |
| YCR032W | BPH1 | unknown | -0.5042 | -1.4 | 0.00446393 |
| YDL125C | HNT1 | hydrolase * | -0.5040 | -1.4 | 0.0293068 |
| YER169W | RPH1 | specific transcriptional repressor | -0.5033 | -1.4 | 0.0804288 |
| YMR086W | | unknown | -0.5010 | -1.4 | 0.117261 |
| YGR031C-A | | | -0.5008 | -1.4 | 0.127507 |
| YOR251C | | thiosulfate sulfurtransferase | -0.4997 | -1.4 | 0.0109806 |
| YPR149W | NCE102 | unknown | -0.4990 | -1.4 | 0.0186356 |
| YPL218W | SAR1 | GTPase | -0.4972 | -1.4 | 0.11647 |
| YNL159C | ASI2 | unknown | -0.4970 | -1.4 | 0.0175164 |
| YPR121W | THI22 | phosphomethylpyrimidine kinase | -0.4957 | -1.4 | 0.112944 |
| YJL041W | NSP1 | structural molecule | -0.4953 | -1.4 | 0.145725 |
| YBR043C | QDR3 | multidrug efflux pump | -0.4938 | -1.4 | 0.0159107 |
| YLR411W | CTR3 | copper uptake transporter | -0.4935 | -1.4 | 0.0517532 |
| YBR027C | | | -0.4935 | -1.4 | 0.0798947 |
| YOR317W | FAA1 | long-chain-fatty-acid-CoA ligase | -0.4932 | -1.4 | 0.0126549 |
| YKL103C | LAP4 | aminopeptidase I | -0.4928 | -1.4 | 0.00656555 |
| YLR305C | STT4 | 1-phosphatidylinositol 4-kinase | -0.4907 | -1.4 | 0.106349 |
| YMR168C | CEP3 | DNA bending * | -0.4903 | -1.4 | 0.034848 |
| YGL047W | ALG13 | N-acetylglucosaminyldiphosphodolichol N-acetylglucosam | -0.4893 | -1.4 | 0.00410846 |
| YOR390W | | unknown | -0.4890 | -1.4 | 0.0191167 |
| YBR143C | SUP45 | translation release factor | -0.4887 | -1.4 | 0.0665569 |
| YBL112C | | unknown | -0.4852 | -1.4 | 0.141422 |
| YPR204W | | DNA helicase | -0.4845 | -1.4 | 0.0994032 |
| YPL259C | APM1 | clathrin binding | -0.4837 | -1.4 | 0.0289903 |
| YPR008W | HAA1 | specific RNA polymerase II transcription factor | -0.4835 | -1.4 | 0.170798 |
| YGR266W | | unknown | -0.4835 | -1.4 | 0.0707942 |
| YJL092W | HPR5 | DNA helicase | -0.4835 | -1.4 | 0.107225 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YOL145C | CTR9 | RNA polymerase II transcription elongation factor | -0.4830 | -1.4 | 0.0954519 |
| YDL174C | DLD1 | D-lactate dehydrogenase | -0.4830 | -1.4 | 0.0514905 |
| YIR003W | | unknown | -0.4827 | -1.4 | 0.013127 |
| YBR126C | TPS1 | alpha,alpha-trehalose-phosphate synthase (UDP-forming) | -0.4822 | -1.4 | 0.100145 |
| YAL053W | FLC2 | unknown | -0.4820 | -1.4 | 0.0185939 |
| YJR159W | SOR1 | L-iditol 2-dehydrogenase | -0.4813 | -1.4 | 0.0415122 |
| YER104W | RTT105 | unknown | -0.4805 | -1.4 | 0.110018 |
| YKL168C | KKQ8 | protein kinase | -0.4800 | -1.4 | 0.0687454 |
| YJL156C | SSY5 | amino acid binding | -0.4797 | -1.4 | 0.0941579 |
| YEL045C | | | -0.4793 | -1.4 | 0.017107 |
| YNL042W | BOP3 | unknown | -0.4790 | -1.4 | 0.218752 |
| YNL329C | PEX6 | ATPase | -0.4762 | -1.4 | 0.0764895 |
| YIL068C | SEC6 | protein binding | -0.4755 | -1.4 | 0.0509458 |
| YGL022W | STT3 | dolichyl-diphosphooligosaccharide-protein glycotransferase | -0.4752 | -1.4 | 0.358289 |
| YDL245C | HXT15 | glucose transporter | -0.4750 | -1.4 | 0.127009 |
| YMR061W | RNA14 | RNA binding | -0.4738 | -1.4 | 0.0183702 |
| YPR064W | | | -0.4732 | -1.4 | 0.0128985 |
| YJL017W | | | -0.4730 | -1.4 | 0.0485984 |
| YJR094W-A | RPL43B | structural constituent of ribosome | -0.4730 | -1.4 | 0.0610702 |
| YLR043C | TRX1 | thiol-disulfide exchange intermediate | -0.4728 | -1.4 | 0.134268 |
| YDR177W | UBC1 | ubiquitin conjugating enzyme | -0.4717 | -1.4 | 0.00314272 |
| YDR452W | PPN1 | endopolyphosphatase | -0.4717 | -1.4 | 0.00551443 |
| YER060W | FCY21 | cytosine-purine permease | -0.4715 | -1.4 | 0.108139 |
| YJL038C | | unknown | -0.4710 | -1.4 | 0.100531 |
| YGL190C | CDC55 | protein phosphatase type 2A | -0.4700 | -1.4 | 0.0940218 |
| YLR430W | SEN1 | ATP-dependent RNA helicase | -0.4695 | -1.4 | 0.121147 |
| YMR165C | PAH1 | phosphatidate phosphatase | -0.4690 | -1.4 | 0.100722 |
| YJR126C | VPS70 | unknown | -0.4682 | -1.4 | 0.0249087 |
| YIL165C | | unknown | -0.4682 | -1.4 | 0.0870425 |
| YGR187C | HGH1 | unknown | -0.4680 | -1.4 | 0.0805572 |
| YMR028W | TAP42 | protein binding | -0.4670 | -1.4 | 0.021618 |
| YKR095W | MLP1 | ribonucleoprotein binding | -0.4658 | -1.4 | 0.179188 |
| YMR253C | | unknown | -0.4647 | -1.4 | 0.0353179 |
| YGL117W | | unknown | -0.4617 | -1.4 | 0.0369442 |
| YDR219C | MFB1 | protein binding | -0.4617 | -1.4 | 0.129003 |
| YOL129W | VPS68 | unknown | -0.4613 | -1.4 | 0.0118574 |
| YGL113W | SLD3 | chromatin binding | -0.4607 | -1.4 | 0.152581 |
| YOR069W | VPS5 | protein transporter | -0.4605 | -1.4 | 0.0382584 |
| YLR181C | VTA1 | unknown | -0.4602 | -1.4 | 0.0282043 |
| YER035W | EDC2 | RNA binding | -0.4595 | -1.4 | 0.0145424 |
| YFR026C | | unknown | -0.4568 | -1.4 | 0.041594 |
| YBR150C | TBS1 | unknown | -0.4567 | -1.4 | 0.0419611 |
| YER044C | ERG28 | protein binding | -0.4560 | -1.4 | 0.0423537 |
| YJL048C | UBX6 | unknown | -0.4552 | -1.4 | 0.154123 |
| YJL109C | UTP10 | snoRNA binding | -0.4538 | -1.4 | 0.139538 |
| YNR055C | HOL1 | transporter | -0.4538 | -1.4 | 0.250811 |
| YKL198C | PTK1 | protein kinase | -0.4535 | -1.4 | 0.150575 |
| YPR030W | CSR2 | unknown | -0.4532 | -1.4 | 0.0712164 |
| YDR504C | SPG3 | unknown | -0.4522 | -1.4 | 0.0907874 |
| YKL171W | | protein kinase | -0.4517 | -1.4 | 0.172418 |
| YDL097C | RPN6 | structural molecule | -0.4488 | -1.4 | 0.55625 |
| YCR049C | | | -0.4487 | -1.4 | 0.0193044 |
| YER152C | | unknown | -0.4477 | -1.4 | 0.129653 |
| YJL111W | CCT7 | unfolded protein binding | -0.4472 | -1.4 | 0.0712587 |
| YDL094C | | | -0.4470 | -1.4 | 0.084454 |
| YMR040W | YET2 | unknown | -0.4468 | -1.4 | 0.00702652 |
| YDR354W | TRP4 | anthranilate phosphoribosyltransferase | -0.4462 | -1.4 | 0.205888 |
| YBR056W | | unknown | -0.4462 | -1.4 | 0.047543 |
| YGR148C | RPL24B | structural constituent of ribosome* | -0.4455 | -1.4 | 0.00613356 |
| YLL044W | | | -0.4455 | -1.4 | 0.130872 |
| YKR099W | BAS1 | RNA polymerase II transcription factor | -0.4448 | -1.4 | 0.23014 |
| YOR002W | ALG6 | transferase , transferring hexosyl groups | -0.4442 | -1.4 | 0.0486861 |
| YHR029C | YHI9 | unknown | -0.4432 | -1.4 | 0.0151874 |
| YBR208C | DUR1.2 | allophanate hydrolase * | -0.4425 | -1.4 | 0.0451659 |
| YLL042C | ATG10 | APG12 conjugating enzyme | -0.4423 | -1.4 | 0.0536761 |
| YGR026W | | unknown | -0.4422 | -1.4 | 0.0962748 |
| YPR160W | GPH1 | glycogen phosphorylase | -0.4403 | -1.4 | 0.129606 |
| YPL226W | NEW1 | ATPase * | -0.4400 | -1.4 | 0.0484021 |
| YBR241C | | unknown | -0.4395 | -1.4 | 0.0628322 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YCL039W | GID7 | unknown | -0.4393 | -1.4 | 0.0120231 |
| YOR257W | CDC31 | structural constituent of cytoskeleton | -0.4393 | -1.4 | 0.0116983 |
| YCRX13W | | | -0.4392 | -1.4 | 0.053636 |
| YPL135W | ISU1 | protein binding | -0.4388 | -1.4 | 0.0137557 |
| YKR106W | | transporter | -0.4387 | -1.4 | 0.1718 |
| YDR094W | | | -0.4385 | -1.4 | 0.13269 |
| YNL066W | SUN4 | glucosidase | -0.4383 | -1.4 | 0.0675842 |
| YNL288W | CAF40 | unknown | -0.4382 | -1.4 | 0.09862 |
| YLR304C | ACO1 | aconitate hydratase | -0.4375 | -1.4 | 0.317796 |
| YIL090W | ICE2 | unknown | -0.4372 | -1.4 | 0.0491991 |
| YCLX06C | | | -0.4372 | -1.4 | 0.131869 |
| YEL032W | MCM3 | chromatin binding | -0.4352 | -1.4 | 0.0664749 |
| YGL121C | GPG1 | signal transducer | -0.4348 | -1.4 | 0.0307259 |
| YDR180W | SCC2 | unknown | -0.4347 | -1.4 | 0.00701108 |
| YHL050C | | helicase | -0.4345 | -1.4 | 0.0142224 |
| YMR011W | HXT2 | glucose transporter | -0.4338 | -1.4 | 0.000672935 |
| YIR026C | YVH1 | protein tyrosine phosphatase * | -0.4335 | -1.4 | 0.0717492 |
| YPL277C | | unknown | -0.4322 | -1.3 | 0.000202168 |
| YNL238W | KEX2 | serine-type endopeptidase | -0.4322 | -1.3 | 0.325006 |
| YOR162C | YRR1 | specific RNA polymerase II transcription factor | -0.4317 | -1.3 | 0.00379771 |
| YPR014C | | | -0.4305 | -1.3 | 0.224391 |
| YER135C | | | -0.4302 | -1.3 | 0.0369493 |
| YGR224W | AZR1 | azole transporter | -0.4298 | -1.3 | 0.0816517 |
| YBR036C | CSG2 | enzyme regulator | -0.4297 | -1.3 | 0.111793 |
| YOR008C | SLG1 | transmembrane receptor | -0.4295 | -1.3 | 0.00608226 |
| YHR146W | CRP1 | DNA binding | -0.4293 | -1.3 | 0.0820957 |
| YPL260W | | unknown | -0.4292 | -1.3 | 0.206985 |
| YLR422W | | unknown | -0.4290 | -1.3 | 0.12454 |
| YHR217C | | | -0.4290 | -1.3 | 0.057322 |
| YLR403W | SFP1 | transcription factor | -0.4280 | -1.3 | 0.0664659 |
| YMR009W | ADI1 | oxidoreductase | -0.4268 | -1.3 | 0.0113855 |
| YER152W-A | | | -0.4263 | -1.3 | 0.476486 |
| YMR136W | GAT2 | transcription factor | -0.4260 | -1.3 | 0.0599794 |
| YPL105C | | unknown | -0.4238 | -1.3 | 0.0450218 |
| YGR030C | POP6 | ribonuclease P * | -0.4227 | -1.3 | 0.123354 |
| YGL229C | SAP4 | protein serine/threonine phosphatase | -0.4227 | -1.3 | 0.210173 |
| YDR435C | PPM1 | C-terminal protein carboxyl methyltransferase | -0.4223 | -1.3 | 0.017329 |
| YDR458C | | unknown | -0.4220 | -1.3 | 0.0496971 |
| YDR135C | YCF1 | bilirubin transporter | -0.4213 | -1.3 | 0.0178295 |
| YOL020W | TAT2 | aromatic amino acid transporter | -0.4203 | -1.3 | 0.0608652 |
| YHR193C | EGD2 | unfolded protein binding | -0.4197 | -1.3 | 0.0131244 |
| YJL019W | MPS3 | unknown | -0.4193 | -1.3 | 0.0193792 |
| YPR119W | CLB2 | cyclin-dependent protein kinase | -0.4193 | -1.3 | 0.0622373 |
| YGL080W | FMP37 | unknown | -0.4182 | -1.3 | 0.238567 |
| YOR118W | | unknown | -0.4178 | -1.3 | 0.0194561 |
| YKR050W | TRK2 | potassium ion transporter | -0.4178 | -1.3 | 0.133554 |
| YDR070C | FMP16 | unknown | -0.4178 | -1.3 | 0.0459416 |
| YDL016C | | | -0.4177 | -1.3 | 0.0336167 |
| YGL125W | MET13 | structural constituent of ribosome* | -0.4148 | -1.3 | 0.0382414 |
| YJR049C | UTR1 | NADH kinase | -0.4148 | -1.3 | 0.108827 |
| YLR212C | TUB4 | structural constituent of cytoskeleton | -0.4133 | -1.3 | 0.102901 |
| YDL019C | OSH2 | oxysterol binding | -0.4132 | -1.3 | 0.317286 |
| YPL241C | CIN2 | unknown | -0.4122 | -1.3 | 0.0694184 |
| YMR205C | PFK2 | 6-phosphofructokinase | -0.4110 | -1.3 | 0.0300246 |
| YGL146C | | unknown | -0.4103 | -1.3 | 0.0423687 |
| YGL056C | SDS23 | unknown | -0.4100 | -1.3 | 0.0889102 |
| YHL032C | GUT1 | glycerol kinase | -0.4097 | -1.3 | 0.0430023 |
| YOR296W | | unknown | -0.4093 | -1.3 | 0.111869 |
| YOR005C | DNL4 | DNA ligase (ATP) | -0.4087 | -1.3 | 0.427535 |
| YPR111W | DBF20 | protein serine/threonine kinase | -0.4085 | -1.3 | 0.216799 |
| YBR047W | FMP23 | unknown | -0.4080 | -1.3 | 0.0758466 |
| YEL020C | | unknown | -0.4075 | -1.3 | 0.0634133 |
| YDR381W | YRA1 | RNA binding | -0.4072 | -1.3 | 0.112742 |
| YKL036C | | | -0.4070 | -1.3 | 0.0473712 |
| YJL021C | | | -0.4067 | -1.3 | 0.0339177 |
| YOR015W | | | -0.4065 | -1.3 | 0.034166 |
| YCL069W | VBA3 | basic amino acid permease | -0.4065 | -1.3 | 0.0939933 |
| YMR193C-A | | | -0.4062 | -1.3 | 0.170668 |
| YLR231C | BNA5 | kynureninase | -0.4062 | -1.3 | 0.0201093 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-------------|
| YHR219W | | unknown | -0.4060 | -1.3 | 0.0114355 |
| YLL019C | KNS1 | protein serine/threonine kinase | -0.4057 | -1.3 | 0.0525125 |
| YNL055C | POR1 | voltage-gated ion-selective channel | -0.4053 | -1.3 | 0.000681334 |
| YDL090C | RAM1 | protein farnesyltransferase | -0.4053 | -1.3 | 0.0163692 |
| YMR294W | JNM1 | structural constituent of cytoskeleton | -0.4033 | -1.3 | 0.0320005 |
| YMR156C | TPP1 | polynucleotide 3'-phosphatase | -0.4027 | -1.3 | 0.0155728 |
| YIL114C | POR2 | voltage-gated ion-selective channel | -0.4022 | -1.3 | 0.118164 |
| YEL035C | UTR5 | unknown | -0.4022 | -1.3 | 0.0931454 |
| YCL033C | | protein-methionine-R-oxide reductase | -0.4013 | -1.3 | 0.0416362 |
| YCR043C | | unknown | -0.4013 | -1.3 | 0.12404 |
| YDR093W | DNF2 | phospholipid-translocating ATPase | -0.4008 | -1.3 | 0.0682648 |
| YML059C | NTE1 | hydrolase * | -0.4005 | -1.3 | 0.0720435 |
| YKR046C | PET10 | unknown | -0.4003 | -1.3 | 0.0351939 |
| YGR178C | PBP1 | unknown | -0.4002 | -1.3 | 0.26864 |
| YIL146C | ECM37 | unknown | -0.3990 | -1.3 | 0.0281309 |
| YIL173W | VTH1 | signal sequence binding | -0.3988 | -1.3 | 0.0046866 |
| YOR063W | RPL3 | structural constituent of ribosome | -0.3987 | -1.3 | 0.0332418 |
| YEL046C | GLY1 | threonine aldolase | -0.3985 | -1.3 | 0.0326248 |
| YLR286C | CTS1 | endochitinase | -0.3975 | -1.3 | 0.0145287 |
| YBL072C | RPS8A | structural constituent of ribosome | -0.3972 | -1.3 | 0.146455 |
| YHR120W | MSH1 | ATP binding | -0.3972 | -1.3 | 0.11787 |
| YNR056C | BIO5 | permease | -0.3970 | -1.3 | 0.201742 |
| YGL026C | TRP5 | tryptophan synthase | -0.3970 | -1.3 | 0.216121 |
| YNL245C | CWC25 | unknown | -0.3958 | -1.3 | 0.100959 |
| YOR386W | PHR1 | deoxyribodipyrimidine photo-lyase | -0.3957 | -1.3 | 0.0923783 |
| YDR221W | GTB1 | alpha-glucosidase | -0.3952 | -1.3 | 0.256941 |
| YGL020C | GET1 | unknown | -0.3945 | -1.3 | 0.0485926 |
| YGL123W | RPS2 | structural constituent of ribosome | -0.3943 | -1.3 | 0.00716124 |
| YGR230W | BNS1 | unknown | -0.3940 | -1.3 | 0.217864 |
| YMR004W | MVP1 | unknown | -0.3938 | -1.3 | 0.0710217 |
| YBR177C | EHT1 | serine hydrolase | -0.3937 | -1.3 | 0.0193988 |
| YNL139C | RLR1 | DNA binding | -0.3923 | -1.3 | 0.212095 |
| YIL086C | | | -0.3920 | -1.3 | 0.126133 |
| YNL020C | ARK1 | protein serine/threonine kinase | -0.3915 | -1.3 | 0.00441007 |
| YOR071C | | transporter | -0.3910 | -1.3 | 0.0207143 |
| YMR306W | FKS3 | 1,3-beta-glucan synthase | -0.3908 | -1.3 | 0.12844 |
| YGL031C | RPL24A | structural constituent of ribosome* | -0.3905 | -1.3 | 0.0210776 |
| YPR051W | MAK3 | peptide alpha-N-acetyltransferase | -0.3892 | -1.3 | 0.01282 |
| YGL134W | PCL10 | cyclin-dependent protein kinase | -0.3890 | -1.3 | 0.0825362 |
| YNL025C | SSN8 | general RNA polymerase II transcription factor | -0.3888 | -1.3 | 0.00675383 |
| YBR074W | | metalloendopeptidase | -0.3883 | -1.3 | 0.0210965 |
| YDR232W | HEM1 | 5-aminolevulinate synthase | -0.3880 | -1.3 | 0.0214868 |
| YJR036C | HUL4 | ubiquitin-protein ligase | -0.3877 | -1.3 | 0.0582429 |
| YCR057C | PWP2 | snoRNA binding | -0.3872 | -1.3 | 0.103727 |
| YNL113W | RPC19 | DNA-directed RNA polymerase | -0.3867 | -1.3 | 0.0122487 |
| YGR110W | | unknown | -0.3867 | -1.3 | 0.0942809 |
| YKR010C | TOF2 | unknown | -0.3865 | -1.3 | 0.0590761 |
| YHL015W | RPS20 | structural constituent of ribosome | -0.3862 | -1.3 | 0.0396134 |
| YJL015C | | | -0.3860 | -1.3 | 0.0625249 |
| YOR206W | NOC2 | unknown | -0.3855 | -1.3 | 0.0966049 |
| YPL019C | VTC3 | unknown | -0.3855 | -1.3 | 0.213552 |
| YHR155W | YSP1 | unknown | -0.3853 | -1.3 | 0.000390112 |
| YOL015W | IRC10 | unknown | -0.3853 | -1.3 | 0.0130453 |
| YBL048W | | | -0.3850 | -1.3 | 0.0300682 |
| YGR044C | RME1 | specific transcriptional repressor | -0.3847 | -1.3 | 0.121294 |
| YCR070W | | | -0.3845 | -1.3 | 0.0157626 |
| YDR130C | FIN1 | unknown | -0.3835 | -1.3 | 0.0467969 |
| YLL024C | SSA2 | ATPase * | -0.3832 | -1.3 | 0.17181 |
| YOL075C | | ATPase , coupled to transmembrane movement of substance | -0.3830 | -1.3 | 0.375379 |
| YDR030C | RAD28 | unknown | -0.3827 | -1.3 | 0.11101 |
| YLR096W | KIN2 | protein kinase | -0.3827 | -1.3 | 0.0345304 |
| YLR101C | | | -0.3817 | -1.3 | 0.211039 |
| YLR149C | | unknown | -0.3807 | -1.3 | 0.0435502 |
| YCR079W | | phosphoprotein phosphatase | -0.3807 | -1.3 | 0.16315 |
| YER064C | | unknown | -0.3805 | -1.3 | 0.123981 |
| YNL218W | MGS1 | ATPase * | -0.3798 | -1.3 | 0.249181 |
| YJR047C | ANB1 | translation initiation factor | -0.3798 | -1.3 | 0.0321636 |
| YDL223C | HBT1 | unknown | -0.3798 | -1.3 | 0.0913413 |
| YGR133W | PEX4 | ubiquitin conjugating enzyme | -0.3797 | -1.3 | 0.0420709 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-------------|
| YMR155W | | unknown | -0.3795 | -1.3 | 0.0297039 |
| YIL151C | | unknown | -0.3793 | -1.3 | 0.0826828 |
| YLL048C | YBT1 | ATPase , coupled to transmembrane movement of substance | -0.3790 | -1.3 | 0.148646 |
| YDR488C | PAC11 | microtubule motor * | -0.3788 | -1.3 | 0.0181579 |
| YLL056C | | unknown | -0.3787 | -1.3 | 0.0986739 |
| YDR255C | RMD5 | unknown | -0.3765 | -1.3 | 0.0107499 |
| YLR046C | | unknown | -0.3762 | -1.3 | 0.0308213 |
| YPR098C | | unknown | -0.3758 | -1.3 | 0.182261 |
| YLR004C | | transporter | -0.3758 | -1.3 | 0.166659 |
| YJL214W | HXT8 | glucose transporter | -0.3758 | -1.3 | 0.0182851 |
| YKL213C | DOA1 | ubiquitin binding | -0.3753 | -1.3 | 0.212179 |
| YLR281C | | unknown | -0.3753 | -1.3 | 0.0953294 |
| YPR112C | MRD1 | snoRNA binding | -0.3752 | -1.3 | 0.332033 |
| YHR053C | CUP1-1 | copper ion binding | -0.3748 | -1.3 | 0.0818693 |
| YDR286C | | unknown | -0.3742 | -1.3 | 0.167057 |
| YFR009W | GCN20 | ATPase | -0.3740 | -1.3 | 0.041979 |
| YJL050W | MTR4 | ATP-dependent RNA helicase | -0.3733 | -1.3 | 0.117821 |
| YKL112W | ABF1 | DNA binding | -0.3732 | -1.3 | 0.140515 |
| YNL311C | SKP2 | protein binding | -0.3730 | -1.3 | 0.162079 |
| YOR109W | INP53 | inositol-polyphosphate 5-phosphatase | -0.3730 | -1.3 | 0.103584 |
| YJR002W | MPP10 | unknown | -0.3725 | -1.3 | 0.0935528 |
| YCL029C | BIK1 | microtubule binding | -0.3725 | -1.3 | 0.149847 |
| YAL017W | PSK1 | protein serine/threonine kinase | -0.3723 | -1.3 | 0.00339556 |
| YEL013W | VAC8 | protein binding | -0.3717 | -1.3 | 0.0590419 |
| YJL044C | GYP6 | GTPase activator | -0.3717 | -1.3 | 0.0957658 |
| YML048W-A | | | -0.3710 | -1.3 | 0.211523 |
| YER110C | KAP123 | protein carrier | -0.3697 | -1.3 | 0.085444 |
| YEL067C | | | -0.3693 | -1.3 | 0.0814103 |
| YBL001C | ECM15 | unknown | -0.3693 | -1.3 | 0.0301314 |
| YEL018C-A | | | -0.3690 | -1.3 | 0.219798 |
| YDR464W | SPP41 | unknown | -0.3690 | -1.3 | 0.0662367 |
| YIL045W | PIG2 | protein phosphatase type 1 regulator | -0.3687 | -1.3 | 0.00277491 |
| YNL271C | BNI1 | cytoskeletal regulator | -0.3683 | -1.3 | 0.0397408 |
| YPL115C | BEM3 | signal transducer * | -0.3682 | -1.3 | 0.26892 |
| YJR084W | CSN12 | unknown | -0.3680 | -1.3 | 0.0573032 |
| YBR221C | PDB1 | pyruvate dehydrogenase | -0.3673 | -1.3 | 0.00295331 |
| YKL131W | | | -0.3672 | -1.3 | 0.309144 |
| YIR010W | DSN1 | unknown | -0.3670 | -1.3 | 0.0920181 |
| YMR071C | TVP18 | unknown | -0.3667 | -1.3 | 0.105491 |
| YPL080C | | | -0.3663 | -1.3 | 0.275041 |
| YJR061W | | unknown | -0.3663 | -1.3 | 0.148606 |
| YDL238C | GUD1 | hydrolase , acting on carbon-nitrogen (but not peptide) bond | -0.3662 | -1.3 | 0.0237509 |
| YJR044C | VPS55 | unknown | -0.3658 | -1.3 | 0.0182088 |
| YER033C | ZRG8 | unknown | -0.3655 | -1.3 | 0.100425 |
| YBR270C | BIT2 | unknown | -0.3655 | -1.3 | 0.0146778 |
| YDR036C | EHD3 | 3-hydroxyisobutyryl-CoA hydrolase | -0.3653 | -1.3 | 0.0225316 |
| YBR084W | MIS1 | formate-tetrahydrofolate ligase * | -0.3652 | -1.3 | 0.118197 |
| YKL002W | DID4 | protein binding | -0.3652 | -1.3 | 0.147969 |
| YLR095C | IOC2 | protein binding | -0.3650 | -1.3 | 0.0790264 |
| YBR049C | REB1 | RNA polymerase II transcription factor | -0.3648 | -1.3 | 0.031966 |
| YLR297W | | unknown | -0.3648 | -1.3 | 0.0477301 |
| YNL099C | OCA1 | protein tyrosine phosphatase | -0.3648 | -1.3 | 0.0157561 |
| YOR302W | | translation regulator | -0.3647 | -1.3 | 0.109012 |
| YGL019W | CKB1 | protein kinase | -0.3645 | -1.3 | 0.223894 |
| YJL027C | | unknown | -0.3643 | -1.3 | 0.000625393 |
| YGL180W | ATG1 | protein serine/threonine kinase | -0.3643 | -1.3 | 0.196206 |
| YCR026C | NPP1 | nucleoside-triphosphatase * | -0.3643 | -1.3 | 0.0910544 |
| YDL043C | PRP11 | RNA binding | -0.3635 | -1.3 | 0.187732 |
| YOL068C | HST1 | NAD-dependent histone deacetylase * | -0.3635 | -1.3 | 0.0669284 |
| YOR377W | ATF1 | alcohol O-acetyltransferase | -0.3633 | -1.3 | 0.0506751 |
| YML115C | VAN1 | mannosyltransferase | -0.3628 | -1.3 | 0.0858051 |
| YNL047C | SLM2 | phosphoinositide binding | -0.3623 | -1.3 | 0.115462 |
| YOR389W | | unknown | -0.3620 | -1.3 | 0.347217 |
| YGR032W | GSC2 | 1,3-beta-glucan synthase | -0.3615 | -1.3 | 0.102087 |
| YGL147C | RPL9A | structural constituent of ribosome | -0.3613 | -1.3 | 0.0266679 |
| YPR027C | | unknown | -0.3612 | -1.3 | 0.0389922 |
| YIL154C | IMP2' | transcription coactivator | -0.3612 | -1.3 | 0.0337666 |
| YER114C | BOI2 | phospholipid binding | -0.3608 | -1.3 | 0.261809 |
| YBR125C | PTC4 | protein phosphatase type 2C | -0.3602 | -1.3 | 0.00616517 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-------------|
| YBL049W | MOH1 | unknown | -0.3602 | -1.3 | 0.00186263 |
| YJL094C | KHA1 | potassium:hydrogen antiporter | -0.3600 | -1.3 | 0.168465 |
| YIR020C | | | -0.3595 | -1.3 | 0.171964 |
| YGL195W | GCN1 | unknown | -0.3593 | -1.3 | 0.2328 |
| YFR040W | SAP155 | protein serine/threonine phosphatase | -0.3588 | -1.3 | 0.200062 |
| YGL149W | | | -0.3587 | -1.3 | 0.0303589 |
| YIL078W | THS1 | threonine-tRNA ligase | -0.3587 | -1.3 | 0.0608405 |
| YLR466W | YRF1-4 | DNA helicase | -0.3580 | -1.3 | 0.230476 |
| YCL001W | RER1 | unknown | -0.3578 | -1.3 | 0.130139 |
| YKR054C | DYN1 | motor | -0.3578 | -1.3 | 0.0243847 |
| YDR114C | | | -0.3573 | -1.3 | 0.25864 |
| YER126C | NSA2 | unknown | -0.3563 | -1.3 | 0.125991 |
| YBR302C | COS2 | unknown | -0.3562 | -1.3 | 0.353745 |
| YJL087C | TRL1 | 2',3'-cyclic-nucleotide 3'-phosphodiesterase * | -0.3557 | -1.3 | 0.15026 |
| YER164W | CHD1 | ATPase * | -0.3547 | -1.3 | 0.339312 |
| YAL064W | | unknown | -0.3547 | -1.3 | 0.0305268 |
| YOR173W | DCS2 | unknown | -0.3542 | -1.3 | 0.181664 |
| YNL103W | MET4 | transcription coactivator | -0.3540 | -1.3 | 0.00229351 |
| YGR270W | YTA7 | ATPase | -0.3533 | -1.3 | 0.0281176 |
| YLR220W | CCC1 | manganese ion transporter | -0.3525 | -1.3 | 0.080686 |
| YKR081C | RPF2 | rRNA binding | -0.3513 | -1.3 | 0.0568543 |
| YAL010C | MDM10 | unknown | -0.3507 | -1.3 | 0.000520101 |
| YNL194C | | unknown | -0.3505 | -1.3 | 0.114887 |
| YLR172C | DPH5 | diphthine synthase | -0.3502 | -1.3 | 0.0386634 |
| YJL034W | KAR2 | ATPase * | -0.3498 | -1.3 | 0.0953108 |
| YKR028W | SAP190 | protein serine/threonine phosphatase | -0.3488 | -1.3 | 0.0233924 |
| YNR062C | | unknown | -0.3485 | -1.3 | 0.11698 |
| YGR268C | HUA1 | unknown | -0.3478 | -1.3 | 0.110137 |
| YIR007W | | unknown | -0.3477 | -1.3 | 0.00633846 |
| YGL045W | RIM8 | unknown | -0.3473 | -1.3 | 0.361064 |
| YMR198W | CIK1 | microtubule motor | -0.3470 | -1.3 | 0.0889176 |
| YJR025C | BNA1 | 3-hydroxyanthranilate 3,4-dioxygenase | -0.3467 | -1.3 | 0.034792 |
| YIL174W | | unknown | -0.3467 | -1.3 | 0.0408922 |
| YLR357W | RSC2 | unknown | -0.3462 | -1.3 | 0.0704888 |
| YOL025W | LAG2 | unknown | -0.3460 | -1.3 | 0.153501 |
| YLR406C | RPL31B | structural constituent of ribosome | -0.3455 | -1.3 | 0.424517 |
| YHR026W | PPA1 | hydrogen-transporting ATPase , rotational mechanism | -0.3453 | -1.3 | 0.000658306 |
| YDR040C | ENA1 | ATPase , coupled to transmembrane movement of ions, phc | -0.3452 | -1.3 | 0.0573736 |
| YLR247C | | helicase | -0.3448 | -1.3 | 0.0604129 |
| YPL087W | YDC1 | ceramidase | -0.3438 | -1.3 | 0.0427593 |
| YGL028C | SCW11 | glucan 1,3-beta-glucosidase | -0.3438 | -1.3 | 0.0161148 |
| YML103C | NUP188 | structural molecule | -0.3433 | -1.3 | 0.0702357 |
| YDR128W | | unknown | -0.3433 | -1.3 | 0.0264648 |
| YBR129C | OPY1 | unknown | -0.3430 | -1.3 | 0.0340172 |
| YEL012W | UBC8 | ubiquitin conjugating enzyme | -0.3430 | -1.3 | 0.032993 |
| YJL190C | RPS22A | structural constituent of ribosome | -0.3427 | -1.3 | 0.0309105 |
| YCR030C | SYP1 | unknown | -0.3425 | -1.3 | 0.00612528 |
| YNL189W | SRP1 | protein carrier | -0.3418 | -1.3 | 0.155534 |
| YPL003W | ULA1 | NEDD8 activating enzyme | -0.3408 | -1.3 | 0.0551799 |
| YML049C | RSE1 | U2 snRNA binding | -0.3407 | -1.3 | 0.117339 |
| YGR033C | TIM21 | unknown | -0.3405 | -1.3 | 0.394495 |
| YPL196W | OXR1 | unknown | -0.3403 | -1.3 | 0.123484 |
| YOR289W | | unknown | -0.3395 | -1.3 | 0.00946276 |
| YFR024C | | | -0.3393 | -1.3 | 0.193063 |
| YKR048C | NAP1 | protein binding | -0.3367 | -1.3 | 0.256456 |
| YGL104C | VPS73 | unknown | -0.3365 | -1.3 | 0.181702 |
| YBL037W | APL3 | unknown | -0.3362 | -1.3 | 0.36038 |
| YCL034W | LSB5 | unknown | -0.3362 | -1.3 | 0.110745 |
| YBR206W | | | -0.3350 | -1.3 | 0.16347 |
| YNL255C | GIS2 | unknown | -0.3345 | -1.3 | 0.113514 |
| YNL273W | TOF1 | unknown | -0.3345 | -1.3 | 0.0420212 |
| YCR061W | | unknown | -0.3342 | -1.3 | 0.0100186 |
| YBR295W | PCA1 | ATPase , coupled to transmembrane movement of ions, phc | -0.3337 | -1.3 | 0.0899295 |
| YER132C | PMD1 | unknown | -0.3333 | -1.3 | 0.0230068 |
| YLR079W | SIC1 | protein binding | -0.3332 | -1.3 | 0.146741 |
| YIL130W | ASG1 | unknown | -0.3330 | -1.3 | 0.0302388 |
| YBL023C | MCM2 | chromatin binding | -0.3330 | -1.3 | 0.357925 |
| YJL150W | | | -0.3328 | -1.3 | 0.338089 |
| YPR022C | | unknown | -0.3323 | -1.3 | 0.010152 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|--|----------------------------|-------------|-------------|
| YDR150W | NUM1 | tubulin binding | -0.3322 | -1.3 | 0.329595 |
| YNL339C | YRF1-6 | DNA helicase | -0.3308 | -1.3 | 0.135011 |
| YBR184W | | unknown | -0.3305 | -1.3 | 0.0735225 |
| YGL096W | TOS8 | transcription factor | -0.3300 | -1.3 | 0.028522 |
| YCR046C | IMG1 | structural constituent of ribosome | -0.3300 | -1.3 | 0.155281 |
| YPR023C | EAF3 | histone acetyltransferase | -0.3288 | -1.3 | 0.253674 |
| YIR033W | MGA2 | transcriptional activator | -0.3287 | -1.3 | 0.252323 |
| YNL064C | YDJ1 | chaperone regulator | -0.3282 | -1.3 | 0.0987389 |
| YGR284C | ERV29 | unknown | -0.3280 | -1.3 | 0.144333 |
| YDR334W | SWR1 | helicase | -0.3277 | -1.3 | 0.338694 |
| YOR253W | NAT5 | peptide alpha-N-acetyltransferase | -0.3273 | -1.3 | 0.376428 |
| YCR024C-A | PMP1 | enzyme regulator | -0.3268 | -1.3 | 0.0436619 |
| YLR441C | RPS1A | structural constituent of ribosome | -0.3268 | -1.3 | 0.000738866 |
| YLL045C | RPL8B | structural constituent of ribosome | -0.3267 | -1.3 | 0.075163 |
| YIR022W | SEC11 | signal peptidase | -0.3267 | -1.3 | 0.0228676 |
| YLR028C | ADE16 | IMP cyclohydrolase * | -0.3265 | -1.3 | 0.731278 |
| YOR221C | MCT1 | [acyl-carrier protein] S-malonyltransferase | -0.3250 | -1.3 | 0.107038 |
| YOR056C | NOB1 | protein binding | -0.3238 | -1.3 | 0.0649157 |
| YOR164C | | unknown | -0.3233 | -1.3 | 0.0459013 |
| YDR267C | CIA1 | unknown | -0.3228 | -1.3 | 0.0736498 |
| YMR148W | | unknown | -0.3222 | -1.3 | 0.146561 |
| YER090W | TRP2 | anthranilate synthase | -0.3222 | -1.3 | 0.239247 |
| YEL077C | | helicase | -0.3222 | -1.3 | 0.0230515 |
| YOR226C | ISU2 | unknown | -0.3222 | -1.3 | 0.26995 |
| YKR058W | GLG1 | glycogenin glucosyltransferase | -0.3218 | -1.2 | 0.00460518 |
| YHR015W | MIP6 | RNA binding | -0.3218 | -1.2 | 0.144614 |
| YKL093W | MBR1 | unknown | -0.3217 | -1.2 | 0.00156233 |
| YLR170C | APS1 | clathrin binding | -0.3213 | -1.2 | 0.137038 |
| YIL121W | QDR2 | multidrug efflux pump | -0.3208 | -1.2 | 0.115464 |
| YCR069W(1) | | | -0.3203 | -1.2 | 0.0464665 |
| YDL087C | LUC7 | mRNA binding | -0.3197 | -1.2 | 0.195616 |
| YJL058C | BIT61 | unknown | -0.3195 | -1.2 | 0.425197 |
| YOL149W | DCP1 | mRNA binding | -0.3192 | -1.2 | 0.0292123 |
| YIL087C | | unknown | -0.3190 | -1.2 | 0.149801 |
| YJL068C | | S-formylglutathione hydrolase | -0.3190 | -1.2 | 0.0641067 |
| YML057W | CMP2 | calcium-dependent protein serine/threonine phosphatase | -0.3190 | -1.2 | 0.286472 |
| YDR304C | CPR5 | peptidyl-prolyl cis-trans isomerase | -0.3187 | -1.2 | 0.167846 |
| YKR066C | CCP1 | cytochrome-c peroxidase | -0.3187 | -1.2 | 0.069421 |
| YLR278C | | unknown | -0.3185 | -1.2 | 0.171956 |
| YMR138W | CIN4 | GTP binding | -0.3177 | -1.2 | 0.203049 |
| YLL055W | | ion transporter | -0.3177 | -1.2 | 0.281899 |
| YJL215C | | | -0.3177 | -1.2 | 0.373016 |
| YGL008C | PMA1 | hydrogen-exporting ATPase , phosphorylative mechanism | -0.3173 | -1.2 | 0.427074 |
| YCL012W | | | -0.3173 | -1.2 | 0.38203 |
| YGL252C | RTG2 | transcription regulator | -0.3172 | -1.2 | 0.253311 |
| YPL039W | | unknown | -0.3168 | -1.2 | 0.190561 |
| YHR213W | | unknown | -0.3163 | -1.2 | 0.127817 |
| YLR013W | GAT3 | transcription factor | -0.3143 | -1.2 | 0.224647 |
| YJL018W | | | -0.3142 | -1.2 | 0.240913 |
| YLR007W | NSE1 | unknown | -0.3133 | -1.2 | 0.0763403 |
| YJR103W | URA8 | CTP synthase | -0.3130 | -1.2 | 0.0488268 |
| YMR026C | PEX12 | protein binding | -0.3127 | -1.2 | 0.222365 |
| YOR307C | SLY41 | unknown | -0.3125 | -1.2 | 0.120842 |
| YJL212C | OPT1 | oligopeptide transporter | -0.3123 | -1.2 | 0.0175854 |
| YJR074W | MOG1 | Ran GTPase binding | -0.3117 | -1.2 | 0.0057921 |
| YOL119C | MCH4 | transporter | -0.3113 | -1.2 | 0.0459738 |
| YBR148W | YSW1 | unknown | -0.3112 | -1.2 | 0.0098082 |
| YPL216W | | unknown | -0.3103 | -1.2 | 0.429185 |
| YGR119C | NUP57 | structural molecule | -0.3103 | -1.2 | 0.0849686 |
| YJL188C | | | -0.3102 | -1.2 | 0.154474 |
| YDL172C | | | -0.3100 | -1.2 | 0.315286 |
| YPR069C | SPE3 | spermidine synthase | -0.3098 | -1.2 | 0.11871 |
| YDL221W | | | -0.3095 | -1.2 | 0.258251 |
| YMR119W | ASI1 | ubiquitin-protein ligase | -0.3095 | -1.2 | 0.0904633 |
| YLR127C | APC2 | protein binding | -0.3095 | -1.2 | 0.00337982 |
| YER115C | SPR6 | unknown | -0.3095 | -1.2 | 0.0457322 |
| YOL115W | PAP2 | polynucleotide adenyltransferase * | -0.3093 | -1.2 | 0.216002 |
| YBR232C | | | -0.3093 | -1.2 | 0.0533928 |
| YNL317W | PFS2 | RNA binding | -0.3088 | -1.2 | 0.241934 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YLR208W | SEC13 | structural molecule | -0.3085 | -1.2 | 0.0143548 |
| YGR117C | | unknown | -0.3085 | -1.2 | 0.0653263 |
| YEL041W | YEF1 | NADH kinase | -0.3083 | -1.2 | 0.0261819 |
| YML105C | SEC65 | unknown | -0.3078 | -1.2 | 0.1293 |
| YDL132W | CDC53 | structural molecule * | -0.3075 | -1.2 | 0.254634 |
| YNL040W | | unknown | -0.3070 | -1.2 | 0.297549 |
| YCL017C | NFS1 | cystathionine gamma-lyase * | -0.3062 | -1.2 | 0.0663081 |
| YPR011C | | transporter | -0.3060 | -1.2 | 0.325417 |
| YNL331C | AAD14 | aryl-alcohol dehydrogenase | -0.3058 | -1.2 | 0.102079 |
| YAL008W | FUN14 | unknown | -0.3048 | -1.2 | 0.0436598 |
| YJL028W | | | -0.3047 | -1.2 | 0.21731 |
| YHR158C | KEL1 | unknown | -0.3045 | -1.2 | 0.0207596 |
| YNL032W | SIW14 | protein tyrosine phosphatase | -0.3043 | -1.2 | 0.185659 |
| YMR244W | | unknown | -0.3032 | -1.2 | 0.191666 |
| YKR064W | | unknown | -0.3030 | -1.2 | 0.0511864 |
| YCL021W | | | -0.3030 | -1.2 | 0.0797907 |
| YMR284W | YKU70 | RNA binding | -0.3027 | -1.2 | 0.107959 |
| YAL063C | FLO9 | mannose binding | -0.3020 | -1.2 | 0.0230551 |
| YDR034C | LYS14 | transcriptional activator | -0.3012 | -1.2 | 0.279172 |
| YMR167W | MLH1 | DNA binding | -0.3012 | -1.2 | 0.363348 |
| YOR116C | RPO31 | DNA-directed RNA polymerase | -0.3010 | -1.2 | 0.216505 |
| YPL279C | | unknown | -0.3008 | -1.2 | 0.172261 |
| YJR033C | RAV1 | unknown | -0.2998 | -1.2 | 0.270695 |
| YDR118W | APC4 | protein binding | -0.2995 | -1.2 | 0.267451 |
| YBR271W | | S-adenosylmethionine-dependent methyltransferase | -0.2987 | -1.2 | 0.218152 |
| YLR036C | | unknown | -0.2987 | -1.2 | 0.141137 |
| YIL088C | AVT7 | transporter | -0.2980 | -1.2 | 0.117921 |
| YCR029C-A | | | -0.2977 | -1.2 | 0.119231 |
| YDR313C | PIB1 | ubiquitin-protein ligase | -0.2973 | -1.2 | 0.144972 |
| YLR188W | MDL1 | ATPase * | -0.2973 | -1.2 | 0.148506 |
| YBR023C | CHS3 | chitin synthase | -0.2972 | -1.2 | 0.240613 |
| YJR143C | PMT4 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.2968 | -1.2 | 0.1268 |
| YLR091W | | unknown | -0.2967 | -1.2 | 0.11262 |
| YHR102W | KIC1 | kinase | -0.2965 | -1.2 | 0.269345 |
| YMR035W | IMP2 | peptidase * | -0.2965 | -1.2 | 0.2329 |
| YGL244W | RTF1 | RNA polymerase II transcription elongation factor | -0.2960 | -1.2 | 0.196771 |
| YHR191C | CTF8 | unknown | -0.2958 | -1.2 | 0.267098 |
| YLR150W | STM1 | DNA binding | -0.2957 | -1.2 | 0.0116584 |
| YMR066W | SOV1 | unknown | -0.2955 | -1.2 | 0.505338 |
| YNL048W | ALG11 | alpha-1,2-mannosyltransferase | -0.2955 | -1.2 | 0.16061 |
| YHR012W | VPS29 | unknown | -0.2953 | -1.2 | 0.288983 |
| YPR095C | SYT1 | ARF guanyl-nucleotide exchange factor | -0.2947 | -1.2 | 0.0952162 |
| YGR244C | LSC2 | succinate-CoA ligase (ADP-forming) | -0.2945 | -1.2 | 0.168096 |
| YJL200C | ACO2 | aconitate hydratase | -0.2942 | -1.2 | 0.204212 |
| YFL057C | AAD16 | aryl-alcohol dehydrogenase | -0.2938 | -1.2 | 0.00211918 |
| YEL075W-A | | | -0.2938 | -1.2 | 0.00463365 |
| YMR104C | YPK2 | protein kinase | -0.2938 | -1.2 | 0.441384 |
| YGR012W | | cysteine synthase | -0.2935 | -1.2 | 0.0704522 |
| YNL140C | | | -0.2933 | -1.2 | 0.00989018 |
| YOR053W | | | -0.2932 | -1.2 | 0.1043 |
| YNL144C | | unknown | -0.2922 | -1.2 | 0.289503 |
| YLR083C | EMP70 | transporter | -0.2922 | -1.2 | 0.0267471 |
| YDR498C | SEC20 | v-SNARE | -0.2922 | -1.2 | 0.0698801 |
| YNL258C | DSL1 | unknown | -0.2915 | -1.2 | 0.153289 |
| YBR038W | CHS2 | chitin synthase | -0.2912 | -1.2 | 0.134516 |
| YKL212W | SAC1 | inositol or phosphatidylinositol phosphatase | -0.2910 | -1.2 | 0.192238 |
| YHR088W | RPF1 | rRNA primary transcript binding | -0.2908 | -1.2 | 0.209599 |
| YKR042W | UTH1 | unknown | -0.2905 | -1.2 | 0.0138391 |
| YDL096C | | | -0.2903 | -1.2 | 0.360037 |
| YEL033W | | | -0.2903 | -1.2 | 0.0126113 |
| YDR191W | HST4 | DNA binding | -0.2897 | -1.2 | 0.0354186 |
| YGR031W | | unknown | -0.2893 | -1.2 | 0.479593 |
| YJL060W | BNA3 | arylformamidase | -0.2892 | -1.2 | 0.105514 |
| YPL114W | | | -0.2885 | -1.2 | 0.126682 |
| YPL283C | YRF1-7 | DNA helicase | -0.2878 | -1.2 | 0.266183 |
| YHL019C | APM2 | clathrin binding | -0.2870 | -1.2 | 0.359133 |
| YDR432W | NPL3 | mRNA binding | -0.2870 | -1.2 | 0.163522 |
| YLR298C | YHC1 | mRNA binding | -0.2860 | -1.2 | 0.275848 |
| YIL075C | RPN2 | endopeptidase * | -0.2855 | -1.2 | 0.0323434 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|----------|-----------|---|----------------------------|-------------|-------------|
| YIL125W | KGD1 | oxoglutarate dehydrogenase | -0.2850 | -1.2 | 0.145096 |
| YNL116W | DMA2 | unknown | -0.2845 | -1.2 | 0.0612135 |
| YLR165C | PUS5 | pseudouridylate synthase | -0.2842 | -1.2 | 0.110067 |
| YPR182W | SMX3 | RNA splicing factor | -0.2837 | -1.2 | 0.126484 |
| YBR240C | THI2 | transcriptional activator | -0.2837 | -1.2 | 0.12723 |
| YJL095W | BCK1 | MAP kinase | -0.2837 | -1.2 | 0.0360289 |
| YGR051C | | | -0.2833 | -1.2 | 0.282056 |
| YHR055C | CUP1-2 | copper ion binding | -0.2828 | -1.2 | 0.145946 |
| YGR272C | | unknown | -0.2827 | -1.2 | 0.160985 |
| YLR133W | CKI1 | choline kinase | -0.2825 | -1.2 | 0.258537 |
| YMR299C | DYN3 | motor | -0.2825 | -1.2 | 0.0732341 |
| YPL002C | SNF8 | protein binding | -0.2822 | -1.2 | 0.142638 |
| YPL248C | GAL4 | transcriptional activator | -0.2817 | -1.2 | 0.148514 |
| YKL188C | PXA2 | ATPase , coupled to transmembrane movement of substance | -0.2815 | -1.2 | 0.130443 |
| YER162C | RAD4 | damaged DNA binding | -0.2812 | -1.2 | 0.0490349 |
| YLR263W | RED1 | protein binding | -0.2810 | -1.2 | 0.26412 |
| YPL255W | BBP1 | structural constituent of cytoskeleton | -0.2805 | -1.2 | 0.390277 |
| YFL056C | AAD6 | aryl-alcohol dehydrogenase | -0.2802 | -1.2 | 0.403613 |
| YKL160W | ELF1 | RNA polymerase II transcription elongation factor | -0.2800 | -1.2 | 0.0373281 |
| YLR062C | | | -0.2797 | -1.2 | 0.164426 |
| YKL179C | COY1 | unknown | -0.2795 | -1.2 | 0.333529 |
| YNR072W | HXT17 | glucose transporter | -0.2795 | -1.2 | 0.254371 |
| YDL247W | MPH2 | carbohydrate transporter | -0.2787 | -1.2 | 0.0749303 |
| YGL114W | | oligopeptide transporter | -0.2782 | -1.2 | 0.0282769 |
| YER097W | | | -0.2777 | -1.2 | 0.206894 |
| YHR143W | DSE2 | glucan 1,3-beta-glucosidase | -0.2775 | -1.2 | 0.0648855 |
| YPL195W | APL5 | protein binding | -0.2763 | -1.2 | 0.20459 |
| YGR046W | | unknown | -0.2760 | -1.2 | 0.0714618 |
| YDR448W | ADA2 | transcription coactivator | -0.2758 | -1.2 | 0.197634 |
| YOR110W | TFC7 | RNA polymerase III transcription factor | -0.2758 | -1.2 | 0.116424 |
| YGR053C | | unknown | -0.2757 | -1.2 | 0.24044 |
| YBR225W | | unknown | -0.2757 | -1.2 | 0.0331226 |
| YNL072W | RNH201 | ribonuclease H | -0.2753 | -1.2 | 0.0304125 |
| YBR019C | GAL10 | UDP-glucose 4-epimerase | -0.2747 | -1.2 | 0.144732 |
| YML089C | | | -0.2742 | -1.2 | 0.0298623 |
| YDL194W | SNF3 | receptor * | -0.2738 | -1.2 | 0.17203 |
| YML085C | TUB1 | structural constituent of cytoskeleton | -0.2737 | -1.2 | 0.198223 |
| YDR477W | SNF1 | AMP-activated protein kinase | -0.2733 | -1.2 | 0.268546 |
| YDR439W | LRS4 | unknown | -0.2732 | -1.2 | 0.29128 |
| YDR233C | RTN1 | unknown | -0.2730 | -1.2 | 0.000750613 |
| YPL033C | | unknown | -0.2730 | -1.2 | 0.153332 |
| YPR047W | MSF1 | phenylalanine-tRNA ligase | -0.2728 | -1.2 | 0.107901 |
| YOL044W | PEX15 | unknown | -0.2723 | -1.2 | 0.14747 |
| YFL048C | EMP47 | unknown | -0.2717 | -1.2 | 0.0820216 |
| YHR075C | PPE1 | structural constituent of ribosome* | -0.2717 | -1.2 | 0.113036 |
| YBL051C | PIN4 | unknown | -0.2713 | -1.2 | 0.127455 |
| YOR197W | MCA1 | caspase | -0.2712 | -1.2 | 0.0943845 |
| YNL007C | SIS1 | unfolded protein binding | -0.2712 | -1.2 | 0.196744 |
| YJL185C | | unknown | -0.2710 | -1.2 | 0.0217942 |
| YBR219C | | unknown | -0.2708 | -1.2 | 0.124366 |
| YBL104C | | unknown | -0.2707 | -1.2 | 0.197818 |
| Q0140(1) | | | -0.2707 | -1.2 | 0.141678 |
| YMR271C | URA10 | orotate phosphoribosyltransferase | -0.2705 | -1.2 | 0.0404899 |
| YLR350W | ORM2 | unknown | -0.2703 | -1.2 | 0.0341071 |
| YFR001W | LOC1 | mRNA binding | -0.2703 | -1.2 | 0.187593 |
| YJL149W | | unknown | -0.2698 | -1.2 | 0.188421 |
| YLR373C | VID22 | unknown | -0.2690 | -1.2 | 0.0736018 |
| YMR143W | RPS16A | structural constituent of ribosome | -0.2688 | -1.2 | 0.000614648 |
| YDR172W | SUP35 | translation release factor | -0.2688 | -1.2 | 0.457449 |
| YGL150C | INO80 | ATPase * | -0.2687 | -1.2 | 0.0351436 |
| YHR079C | IRE1 | protein serine/threonine kinase | -0.2687 | -1.2 | 0.444175 |
| YIL054W | | | -0.2677 | -1.2 | 0.0593568 |
| YNR013C | PHO91 | phosphate transporter | -0.2675 | -1.2 | 0.432888 |
| YER144C | UBP5 | ubiquitin-specific protease | -0.2675 | -1.2 | 0.0653402 |
| YBR123C | TFC1 | RNA polymerase III transcription factor | -0.2670 | -1.2 | 0.0248508 |
| YLR399C | BDF1 | transcription regulator | -0.2670 | -1.2 | 0.00886552 |
| YML086C | ALO1 | D-arabinono-1,4-lactone oxidase | -0.2668 | -1.2 | 0.427595 |
| YPR024W | YME1 | ATP-dependent peptidase | -0.2667 | -1.2 | 0.0419777 |
| YOR018W | ROD1 | unknown | -0.2667 | -1.2 | 0.0374907 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|------------|
| YKL026C | GPX1 | glutathione peroxidase | -0.2658 | -1.2 | 0.0373404 |
| YJR146W | | | -0.2655 | -1.2 | 0.0868322 |
| YPL084W | BRO1 | intracellular transporter | -0.2650 | -1.2 | 0.0688557 |
| YBL079W | NUP170 | structural molecule | -0.2645 | -1.2 | 0.115582 |
| YOR291W | | unknown | -0.2645 | -1.2 | 0.273591 |
| YEL017W | GTT3 | unknown | -0.2643 | -1.2 | 0.0220221 |
| YIL108W | | metalloendopeptidase | -0.2642 | -1.2 | 0.324628 |
| YPR185W | ATG13 | protein binding | -0.2638 | -1.2 | 0.313024 |
| YBR180W | DTR1 | multidrug transporter | -0.2632 | -1.2 | 0.104528 |
| YDR436W | PPZ2 | protein serine/threonine phosphatase | -0.2632 | -1.2 | 0.126736 |
| YNL237W | YTP1 | unknown | -0.2632 | -1.2 | 0.250947 |
| YLR378C | SEC61 | protein transporter | -0.2630 | -1.2 | 0.0174894 |
| YJR135C | MCM22 | protein binding | -0.2627 | -1.2 | 0.107988 |
| YLR464W | | unknown | -0.2625 | -1.2 | 0.0894882 |
| YCL052C | PBN1 | mannosyltransferase | -0.2623 | -1.2 | 0.0618867 |
| YDR157W | | | -0.2620 | -1.2 | 0.2262 |
| YGL241W | KAP114 | protein carrier | -0.2618 | -1.2 | 0.217246 |
| YKL015W | PUT3 | specific RNA polymerase II transcription factor | -0.2618 | -1.2 | 0.0342445 |
| YJL176C | SWI3 | general RNA polymerase II transcription factor | -0.2618 | -1.2 | 0.0275564 |
| YLR129W | DIP2 | snoRNA binding | -0.2615 | -1.2 | 0.289882 |
| YDL184C | RPL41A | structural constituent of ribosome | -0.2610 | -1.2 | 0.314721 |
| YPL222W | FMP40 | unknown | -0.2608 | -1.2 | 0.0572508 |
| YER151C | UBP3 | ubiquitin-specific protease | -0.2608 | -1.2 | 0.0833568 |
| YML019W | OST6 | dolichyl-diphosphooligosaccharide-protein glycotransferase | -0.2603 | -1.2 | 0.0412345 |
| YJR123W | RPS5 | structural constituent of ribosome | -0.2598 | -1.2 | 0.0326675 |
| YLL018C | DPS1 | RNA binding | -0.2595 | -1.2 | 0.0149096 |
| YLR449W | FPR4 | peptidyl-prolyl cis-trans isomerase | -0.2592 | -1.2 | 0.265286 |
| YCR063W | BUD31 | unknown | -0.2590 | -1.2 | 0.0322333 |
| YDL116W | NUP84 | structural molecule | -0.2588 | -1.2 | 0.488018 |
| YEL034W | HYP2 | protein binding | -0.2587 | -1.2 | 0.0214605 |
| YDR390C | UBA2 | SUMO activating enzyme | -0.2587 | -1.2 | 0.107489 |
| YCR075C | ERS1 | L-cystine transporter | -0.2583 | -1.2 | 0.345851 |
| YNL297C | MON2 | guanyl-nucleotide exchange factor | -0.2573 | -1.2 | 0.158648 |
| YAL022C | FUN26 | nucleoside transporter | -0.2573 | -1.2 | 0.154617 |
| YMR008C | PLB1 | lysophospholipase | -0.2567 | -1.2 | 0.128139 |
| YNR019W | ARE2 | sterol O-acyltransferase | -0.2567 | -1.2 | 0.0135564 |
| YDL027C | | unknown | -0.2565 | -1.2 | 0.182748 |
| YMR279C | | unknown | -0.2563 | -1.2 | 0.376012 |
| YGL084C | GUP1 | O-acyltransferase | -0.2560 | -1.2 | 0.00135456 |
| YJL069C | UTP18 | unknown | -0.2560 | -1.2 | 0.0607718 |
| YIL099W | SGA1 | glucan 1,4-alpha-glucosidase | -0.2558 | -1.2 | 0.166555 |
| YGR122W | | unknown | -0.2553 | -1.2 | 0.232435 |
| YML063W | RPS1B | structural constituent of ribosome | -0.2550 | -1.2 | 0.00160181 |
| YJL020C | BBC1 | myosin I binding | -0.2547 | -1.2 | 0.15365 |
| YOL016C | CMK2 | calcium- and calmodulin-dependent protein kinase | -0.2547 | -1.2 | 0.00048993 |
| YLR394W | CST9 | DNA binding | -0.2547 | -1.2 | 0.68103 |
| YBR127C | VMA2 | hydrogen-transporting ATPase , rotational mechanism | -0.2543 | -1.2 | 0.0112857 |
| YHR150W | PEX28 | unknown | -0.2535 | -1.2 | 0.14774 |
| YLR213C | CRR1 | hydrolase , acting on carbon-nitrogen (but not peptide) bonds | -0.2533 | -1.2 | 0.0455935 |
| YLL028W | TPO1 | spermine transporter | -0.2532 | -1.2 | 0.271141 |
| YIL126W | STH1 | ATPase * | -0.2532 | -1.2 | 0.101753 |
| YPL045W | VPS16 | unknown | -0.2530 | -1.2 | 0.0211698 |
| YLR222C | UTP13 | snoRNA binding | -0.2528 | -1.2 | 0.193564 |
| YLR324W | PEX30 | unknown | -0.2523 | -1.2 | 0.489549 |
| YML128C | MSC1 | unknown | -0.2523 | -1.2 | 0.266155 |
| YJL101C | GSH1 | glutamate-cysteine ligase | -0.2522 | -1.2 | 0.00156114 |
| YDR195W | REF2 | RNA binding | -0.2522 | -1.2 | 0.173436 |
| YLR102C | APC9 | protein binding | -0.2520 | -1.2 | 0.103975 |
| YKL104C | GFA1 | glutamine-fructose-6-phosphate transaminase (isomerizing) | -0.2520 | -1.2 | 0.288859 |
| YOR380W | RDR1 | transcription factor | -0.2512 | -1.2 | 0.11804 |
| YOL081W | IRA2 | Ras GTPase activator | -0.2512 | -1.2 | 0.00885719 |
| YKR057W | RPS21A | structural constituent of ribosome | -0.2510 | -1.2 | 0.171322 |
| YMR153W | NUP53 | structural molecule | -0.2502 | -1.2 | 0.367126 |
| YML001W | YPT7 | GTPase | -0.2500 | -1.2 | 0.121734 |
| YLL026W | HSP104 | chaperone binding | -0.2500 | -1.2 | 0.258573 |
| YGR209C | TRX2 | thiol-disulfide exchange intermediate | -0.2495 | -1.2 | 0.154526 |
| YOR152C | | unknown | -0.2495 | -1.2 | 0.220141 |
| YOR104W | PIN2 | unknown | -0.2495 | -1.2 | 0.00179944 |
| YMR015C | ERG5 | C-22 sterol desaturase | -0.2493 | -1.2 | 0.0319063 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|--|----------------------------|-------------|------------|
| YKL215C | | unknown | -0.2492 | -1.2 | 0.41864 |
| YEL044W | IES6 | unknown | -0.2490 | -1.2 | 0.0626487 |
| YLR076C | | | -0.2490 | -1.2 | 0.16775 |
| YAR009C | | | -0.2485 | -1.2 | 0.0969267 |
| YDR424C | DYN2 | microtubule motor | -0.2483 | -1.2 | 0.219492 |
| YMR047C | NUP116 | structural molecule | -0.2482 | -1.2 | 0.520856 |
| YIL107C | PFK26 | 6-phosphofructo-2-kinase | -0.2482 | -1.2 | 0.0170721 |
| YIL164C | NIT1 | nitrilase | -0.2478 | -1.2 | 0.261048 |
| YIL171W | | unknown | -0.2475 | -1.2 | 0.120696 |
| YJL131C | | unknown | -0.2475 | -1.2 | 0.0212596 |
| YDR248C | | unknown | -0.2472 | -1.2 | 0.232051 |
| YLR340W | RPP0 | structural constituent of ribosome | -0.2468 | -1.2 | 0.00877985 |
| YHR171W | ATG7 | APG12 activating enzyme * | -0.2468 | -1.2 | 0.161572 |
| YCRX07W | | | -0.2468 | -1.2 | 0.0810165 |
| YIL002C | INP51 | inositol-polyphosphate 5-phosphatase | -0.2455 | -1.2 | 0.171141 |
| YDL234C | GYP7 | Rab GTPase activator | -0.2455 | -1.2 | 0.057663 |
| YPL050C | MNN9 | mannosyltransferase | -0.2452 | -1.2 | 0.0274268 |
| YPL042C | SSN3 | general RNA polymerase II transcription factor | -0.2450 | -1.2 | 0.0493595 |
| YMR140W | SIP5 | unknown | -0.2448 | -1.2 | 0.273744 |
| YKL076C | | | -0.2442 | -1.2 | 0.0465207 |
| YBR104W | YMC2 | transporter | -0.2438 | -1.2 | 0.0904243 |
| YHL025W | SNF6 | general RNA polymerase II transcription factor | -0.2438 | -1.2 | 0.331243 |
| YDR075W | PPH3 | protein phosphatase type 2A | -0.2437 | -1.2 | 0.15949 |
| YBR204C | | serine hydrolase | -0.2435 | -1.2 | 0.0679402 |
| YMR181C | | unknown | -0.2435 | -1.2 | 0.44701 |
| YML031W | NDC1 | structural constituent of cytoskeleton | -0.2433 | -1.2 | 0.0654127 |
| YJL019W(1) | | | -0.2427 | -1.2 | 0.18281 |
| YBR138C | | unknown | -0.2425 | -1.2 | 0.0809029 |
| YDL203C | ACK1 | unknown | -0.2420 | -1.2 | 0.363991 |
| YDR254W | CHL4 | DNA binding | -0.2417 | -1.2 | 0.126545 |
| YOR282W | | | -0.2410 | -1.2 | 0.0744311 |
| YOR001W | RRP6 | 3'-5'-exoribonuclease | -0.2410 | -1.2 | 0.260877 |
| YHL009W-B | | | -0.2407 | -1.2 | 0.25337 |
| YJL029C | VPS53 | protein binding | -0.2405 | -1.2 | 0.0826482 |
| YFR031C-A | RPL2A | structural constituent of ribosome | -0.2405 | -1.2 | 0.130493 |
| YLR169W | | | -0.2402 | -1.2 | 0.0600343 |
| YLR029C | RPL15A | structural constituent of ribosome* | -0.2402 | -1.2 | 0.153211 |
| YCRX08W | | | -0.2402 | -1.2 | 0.019306 |
| YLR463C | | | -0.2400 | -1.2 | 0.469446 |
| YLR080W | EMP46 | glycoprotein binding | -0.2395 | -1.2 | 0.0869459 |
| YOR329C | SCD5 | protein binding | -0.2395 | -1.2 | 0.300737 |
| YML052W | SUR7 | unknown | -0.2392 | -1.2 | 0.178489 |
| YDR010C | | | -0.2387 | -1.2 | 0.35756 |
| YMR236W | TAF9 | general RNA polymerase II transcription factor | -0.2383 | -1.2 | 0.261216 |
| YMR233W | | unknown | -0.2383 | -1.2 | 0.0724573 |
| YNL061W | NOP2 | S-adenosylmethionine-dependent methyltransferase * | -0.2380 | -1.2 | 0.0521716 |
| YIL167W | SDL1 | L-serine ammonia-lyase | -0.2378 | -1.2 | 0.152492 |
| YLR336C | SGD1 | unknown | -0.2373 | -1.2 | 0.367347 |
| YPL161C | BEM4 | unknown | -0.2373 | -1.2 | 0.0437026 |
| YDR308C | SRB7 | RNA polymerase II transcription mediator | -0.2370 | -1.2 | 0.0643722 |
| YDR205W | MSC2 | cation:cation antiporter | -0.2367 | -1.2 | 0.16205 |
| YOL155C | | glucosidase | -0.2367 | -1.2 | 0.262301 |
| YGR193C | PDX1 | protein binding | -0.2365 | -1.2 | 0.0026172 |
| YJL123C | | unknown | -0.2363 | -1.2 | 0.0973195 |
| YJL204C | RCY1 | protein binding | -0.2363 | -1.2 | 0.144429 |
| YDL075W | RPL31A | structural constituent of ribosome | -0.2360 | -1.2 | 0.197413 |
| YDR159W | SAC3 | protein binding | -0.2355 | -1.2 | 0.214276 |
| YFR014C | CMK1 | calmodulin-dependent protein kinase | -0.2355 | -1.2 | 0.0162826 |
| YDR023W | SES1 | serine-tRNA ligase | -0.2355 | -1.2 | 0.0630756 |
| YDR411C | DFM1 | unknown | -0.2353 | -1.2 | 0.445655 |
| YNL091W | NST1 | unknown | -0.2352 | -1.2 | 0.101224 |
| YAL029C | MYO4 | microfilament motor | -0.2347 | -1.2 | 0.442965 |
| YJR003C | | unknown | -0.2347 | -1.2 | 0.0628078 |
| YGR143W | SKN1 | glucosidase | -0.2343 | -1.2 | 0.511969 |
| YER141W | COX15 | oxidoreductase | -0.2343 | -1.2 | 0.369236 |
| YAL013W | DEP1 | transcription regulator | -0.2342 | -1.2 | 0.0499822 |
| YER056C | FCY2 | cytosine-purine permease | -0.2340 | -1.2 | 0.192632 |
| YCRX11W | | | -0.2338 | -1.2 | 0.147816 |
| YNL293W | MSB3 | Rab GTPase activator | -0.2338 | -1.2 | 0.099886 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YHR126C | | unknown | -0.2337 | -1.2 | 0.045413 |
| YMR027W | | unknown | -0.2337 | -1.2 | 0.153094 |
| YJR124C | | unknown | -0.2332 | -1.2 | 0.320303 |
| YDR007W | TRP1 | phosphoribosylanthranilate isomerase | -0.2330 | -1.2 | 0.249427 |
| YPL243W | SRP68 | signal sequence binding | -0.2328 | -1.2 | 0.314728 |
| YDR257C | SET7 | unknown | -0.2318 | -1.2 | 0.207816 |
| YGR125W | | unknown | -0.2315 | -1.2 | 0.119398 |
| YPR156C | TPO3 | spermine transporter | -0.2310 | -1.2 | 0.322995 |
| YDL140C | RPO21 | DNA-directed RNA polymerase | -0.2310 | -1.2 | 0.19293 |
| YKL028W | TFA1 | general RNA polymerase II transcription factor | -0.2308 | -1.2 | 0.403845 |
| YPL257W | | unknown | -0.2308 | -1.2 | 0.0519107 |
| YKL053W | | | -0.2307 | -1.2 | 0.0269436 |
| YHR050W | SMF2 | di-, tri-valent inorganic cation transporter | -0.2305 | -1.2 | 0.0986259 |
| YER060W-A | FCY22 | cytosine-purine permease | -0.2305 | -1.2 | 0.24289 |
| YCR107W | AAD3 | aryl-alcohol dehydrogenase | -0.2302 | -1.2 | 0.164833 |
| YER176W | ECM32 | DNA helicase | -0.2295 | -1.2 | 0.0259928 |
| YNL323W | LEM3 | phospholipid-translocating ATPase | -0.2295 | -1.2 | 0.0200659 |
| YCR085W | | | -0.2293 | -1.2 | 0.299004 |
| YOR151C | RPB2 | DNA-directed RNA polymerase | -0.2292 | -1.2 | 0.07438 |
| YDR229W | IVY1 | phospholipid binding | -0.2292 | -1.2 | 0.0379493 |
| YOL087C | | unknown | -0.2292 | -1.2 | 0.119349 |
| YLR112W | | | -0.2288 | -1.2 | 0.142866 |
| YDR491C | | | -0.2287 | -1.2 | 0.261307 |
| YDL095W | PMT1 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.2285 | -1.2 | 0.00453395 |
| YNL267W | PIK1 | 1-phosphatidylinositol 4-kinase | -0.2282 | -1.2 | 0.264012 |
| YJL117W | PHO86 | unknown | -0.2278 | -1.2 | 0.0356406 |
| YER029C | SMB1 | RNA splicing factor | -0.2278 | -1.2 | 0.13411 |
| YBL063W | KIP1 | structural constituent of cytoskeleton* | -0.2275 | -1.2 | 0.0721784 |
| YMR283C | RIT1 | transferase , transferring pentosyl groups | -0.2273 | -1.2 | 0.161925 |
| YGL005C | COG7 | unknown | -0.2270 | -1.2 | 0.359984 |
| YDL131W | LYS21 | homocitrate synthase | -0.2268 | -1.2 | 0.0523713 |
| YPL245W | | unknown | -0.2267 | -1.2 | 0.0454598 |
| YAL037W | | unknown | -0.2257 | -1.2 | 0.0056484 |
| YLR187W | SKG3 | unknown | -0.2255 | -1.2 | 0.00387435 |
| YNL041C | COG6 | unknown | -0.2255 | -1.2 | 0.249376 |
| YNL305C | | unknown | -0.2242 | -1.2 | 0.110952 |
| YBR121C | GRS1 | glycine-tRNA ligase | -0.2242 | -1.2 | 0.0784277 |
| YNL216W | RAP1 | DNA binding | -0.2238 | -1.2 | 0.582078 |
| YNR008W | LRO1 | phospholipid:diacylglycerol acyltransferase | -0.2238 | -1.2 | 0.306257 |
| YDR141C | DOP1 | unknown | -0.2237 | -1.2 | 0.11779 |
| YLR022C | SDO1 | unknown | -0.2235 | -1.2 | 0.0481293 |
| YOR324C | FRT1 | unknown | -0.2235 | -1.2 | 0.276188 |
| YPR163C | TIF3 | translation initiation factor | -0.2228 | -1.2 | 0.374347 |
| YBL073W | | | -0.2227 | -1.2 | 0.136236 |
| YLL067C | | helicase | -0.2225 | -1.2 | 0.260464 |
| YDR154C | | | -0.2223 | -1.2 | 0.443918 |
| YLR386W | VAC14 | enzyme activator | -0.2218 | -1.2 | 0.670444 |
| YNL223W | ATG4 | microtubule binding | -0.2215 | -1.2 | 0.177088 |
| YNL249C | MPA43 | unknown | -0.2213 | -1.2 | 0.247201 |
| YDR245W | MNN10 | alpha-1,6-mannosyltransferase | -0.2213 | -1.2 | 0.265929 |
| YIL128W | MET18 | RNA polymerase II transcription factor | -0.2213 | -1.2 | 0.286136 |
| YAL058W | CNE1 | unfolded protein binding | -0.2210 | -1.2 | 0.345723 |
| YDR155C | CPR1 | peptidyl-prolyl cis-trans isomerase | -0.2197 | -1.2 | 0.272882 |
| YDR107C | | unknown | -0.2197 | -1.2 | 0.225775 |
| YER117W | RPL23B | structural constituent of ribosome | -0.2195 | -1.2 | 0.0581939 |
| YPL025C | | | -0.2192 | -1.2 | 0.130259 |
| YHR192W | | unknown | -0.2187 | -1.2 | 0.190976 |
| YJL052W | TDH1 | glyceraldehyde-3-phosphate dehydrogenase | -0.2185 | -1.2 | 0.702105 |
| YFL018C | LPD1 | glycine dehydrogenase | -0.2185 | -1.2 | 0.147204 |
| YNL102W | POL1 | alpha DNA polymerase | -0.2183 | -1.2 | 0.0300448 |
| YBR256C | RIB5 | riboflavin synthase | -0.2182 | -1.2 | 0.159044 |
| YGL016W | KAP122 | protein carrier | -0.2180 | -1.2 | 0.259761 |
| YPL201C | YIG1 | unknown | -0.2177 | -1.2 | 0.15187 |
| YOR044W | | unknown | -0.2175 | -1.2 | 0.481068 |
| YJR045C | SSC1 | ATPase * | -0.2175 | -1.2 | 0.276815 |
| YJL181W | | unknown | -0.2167 | -1.2 | 0.133913 |
| YJL083W | TAX4 | unknown | -0.2165 | -1.2 | 0.11638 |
| YIR017C | MET28 | DNA binding | -0.2163 | -1.2 | 0.0207469 |
| YDL150W | RPC53 | DNA-directed RNA polymerase | -0.2163 | -1.2 | 0.150158 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-------------|
| YNL056W | OCA2 | unknown | -0.2162 | -1.2 | 0.0942471 |
| YCR051W | | unknown | -0.2160 | -1.2 | 0.403227 |
| YGL257C | MNT2 | alpha-1,3-mannosyltransferase | -0.2155 | -1.2 | 0.0547978 |
| YPL200W | CSM4 | unknown | -0.2153 | -1.2 | 0.39695 |
| YOL125W | | unknown | -0.2153 | -1.2 | 0.405998 |
| YPL262W | FUM1 | fumarate hydratase | -0.2152 | -1.2 | 0.236443 |
| YGR231C | PHB2 | unknown | -0.2148 | -1.2 | 0.00187986 |
| YMR092C | AIP1 | protein binding | -0.2147 | -1.2 | 0.292137 |
| YPL055C | LGE1 | unknown | -0.2143 | -1.2 | 0.470635 |
| YML058W | SML1 | enzyme inhibitor | -0.2142 | -1.2 | 0.0839627 |
| YKR027W | BCH2 | unknown | -0.2140 | -1.2 | 0.11813 |
| YDL178W | DLD2 | actin binding | -0.2132 | -1.2 | 0.267848 |
| YIL159W | BNR1 | cytoskeletal protein binding | -0.2132 | -1.2 | 0.0804846 |
| YJL113W | | | -0.2130 | -1.2 | 0.0520398 |
| YPR017C | DSS4 | guanyl-nucleotide exchange factor | -0.2128 | -1.2 | 0.344965 |
| YPL100W | ATG21 | phosphoinositide binding | -0.2118 | -1.2 | 0.331224 |
| YGL142C | GPI10 | mannosyltransferase | -0.2118 | -1.2 | 0.182593 |
| YNL162W | RPL42A | structural constituent of ribosome | -0.2113 | -1.2 | 0.168983 |
| YPL120W | VPS30 | unknown | -0.2110 | -1.2 | 0.46631 |
| YBR223C | TDP1 | tyrosyl-DNA phosphodiesterase | -0.2108 | -1.2 | 0.227007 |
| YPL249C | GYP5 | Rab GTPase activator | -0.2107 | -1.2 | 0.0873048 |
| YER155C | BEM2 | signal transducer * | -0.2105 | -1.2 | 0.385583 |
| YKR018C | | unknown | -0.2105 | -1.2 | 0.403198 |
| YOL093W | TRM10 | tRNA (guanine) methyltransferase | -0.2103 | -1.2 | 0.148813 |
| YGL065C | ALG2 | glycolipid mannosyltransferase | -0.2102 | -1.2 | 0.483334 |
| YKL075C | | unknown | -0.2102 | -1.2 | 0.214572 |
| YLR018C | POM34 | unknown | -0.2098 | -1.2 | 0.302181 |
| YGL208W | SIP2 | AMP-activated protein kinase | -0.2098 | -1.2 | 0.155246 |
| YER069W | ARG5,6 | N-acetyl-gamma-glutamyl-phosphate reductase * | -0.2097 | -1.2 | 0.153587 |
| YDR198C | RKM2 | unknown | -0.2092 | -1.2 | 0.352641 |
| YPL263C | KEL3 | unknown | -0.2083 | -1.2 | 0.189924 |
| YDR239C | | unknown | -0.2082 | -1.2 | 0.0510936 |
| YDR317W | HIM1 | unknown | -0.2080 | -1.2 | 0.131023 |
| YLR353W | BUD8 | unknown | -0.2072 | -1.2 | 0.30901 |
| YOR256C | TRE2 | unknown | -0.2070 | -1.2 | 0.330804 |
| YJL023C | PET130 | unknown | -0.2067 | -1.2 | 0.0228948 |
| YFL042C | | unknown | -0.2060 | -1.2 | 0.223268 |
| YPR031W | NTO1 | unknown | -0.2055 | -1.2 | 0.341158 |
| YNL012W | SPO1 | phospholipase | -0.2050 | -1.2 | 0.282462 |
| YER175C | TMT1 | trans-aconitate 3-methyltransferase | -0.2050 | -1.2 | 0.353242 |
| YHL020C | OPI1 | transcription corepressor | -0.2048 | -1.2 | 0.197546 |
| YMR302C | YME2 | exonuclease | -0.2048 | -1.2 | 0.0712404 |
| YMR121C | RPL15B | structural constituent of ribosome* | -0.2043 | -1.2 | 0.0988629 |
| YML033W | | | -0.2040 | -1.2 | 0.154901 |
| YBR269C | FMP21 | unknown | -0.2040 | -1.2 | 0.0967279 |
| YCR052W | RSC6 | unknown | -0.2038 | -1.2 | 0.145826 |
| YOR209C | NPT1 | nicotinate phosphoribosyltransferase | -0.2037 | -1.2 | 0.523555 |
| YDR022C | CIS1 | unknown | -0.2033 | -1.2 | 0.0556651 |
| YIL113W | SDP1 | MAP kinase | -0.2028 | -1.2 | 0.000606035 |
| YCR106W | RDS1 | DNA binding | -0.2027 | -1.2 | 0.000620939 |
| YNL290W | RFC3 | ATPase * | -0.2023 | -1.2 | 0.069327 |
| YLR384C | IKI3 | RNA polymerase II transcription elongation factor | -0.2022 | -1.2 | 0.222151 |
| YAR029W | | unknown | -0.2022 | -1.2 | 0.51015 |
| YJR137C | ECM17 | sulfite reductase (NADPH) | -0.2018 | -1.2 | 0.271362 |
| YHR164C | DNA2 | ATP-dependent DNA helicase | -0.2018 | -1.2 | 0.127563 |
| YNR007C | ATG3 | APG8 conjugating enzyme | -0.2017 | -1.2 | 0.279633 |
| YOL073C | | unknown | -0.2013 | -1.1 | 0.073663 |
| YLR431C | ATG23 | unknown | -0.2012 | -1.1 | 0.527273 |
| YPR026W | ATH1 | alpha,alpha-trehalase | -0.2010 | -1.1 | 0.434043 |
| YDR533C | HSP31 | unfolded protein binding | -0.2007 | -1.1 | 0.433214 |
| YBR198C | TAF5 | general RNA polymerase II transcription factor | -0.2007 | -1.1 | 0.0946451 |
| YNR047W | | protein kinase | -0.2005 | -1.1 | 0.232978 |
| YLL041C | SDH2 | succinate dehydrogenase | -0.2003 | -1.1 | 0.0204958 |
| YBR200W | BEM1 | protein binding | -0.2003 | -1.1 | 0.355777 |
| YHR124W | NDT80 | transcription factor | -0.2002 | -1.1 | 0.561575 |
| YIL191W | RPS14B | structural constituent of ribosome* | -0.1992 | -1.1 | 0.0196037 |
| YFL004W | VTC2 | unknown | -0.1990 | -1.1 | 0.131676 |
| YIL030C | SSM4 | ubiquitin-protein ligase | -0.1990 | -1.1 | 0.0311752 |
| YIL042C | | kinase | -0.1988 | -1.1 | 0.624707 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YNR071C | | unknown | -0.1987 | -1.1 | 0.014146 |
| YJL055W | | unknown | -0.1983 | -1.1 | 0.218068 |
| YDR103W | STE5 | MAP-kinase | -0.1982 | -1.1 | 0.0622573 |
| YNL182C | IPI3 | unknown | -0.1982 | -1.1 | 0.327416 |
| YLR367W | RPS22B | structural constituent of ribosome | -0.1980 | -1.1 | 0.304604 |
| YNL086W | | unknown | -0.1978 | -1.1 | 0.414778 |
| YAR044W | | | -0.1978 | -1.1 | 0.181802 |
| YJL153W | RRD1 | protein phosphatase type 2A regulator | -0.1977 | -1.1 | 0.446439 |
| YLR413W | | unknown | -0.1973 | -1.1 | 0.14924 |
| YGL077C | HNM1 | choline transporter | -0.1972 | -1.1 | 0.346969 |
| YLR056W | ERG3 | C-5 sterol desaturase | -0.1972 | -1.1 | 0.110014 |
| YOR185C | GSP2 | GTPase | -0.1972 | -1.1 | 0.00874084 |
| YDL083C | RPS16B | structural constituent of ribosome | -0.1968 | -1.1 | 0.18014 |
| YNL111C | CYB5 | electron carrier | -0.1965 | -1.1 | 0.0125685 |
| YER102W | RPS8B | structural constituent of ribosome | -0.1963 | -1.1 | 0.0685175 |
| YGR024C | THG1 | tRNA guanylyltransferase | -0.1963 | -1.1 | 0.0844974 |
| YGL140C | | unknown | -0.1960 | -1.1 | 0.677868 |
| YCR089W | FIG2 | unknown | -0.1960 | -1.1 | 0.202286 |
| YDR063W | | unknown | -0.1957 | -1.1 | 0.43149 |
| YDR514C | | unknown | -0.1955 | -1.1 | 0.00173328 |
| YGR157W | CHO2 | phosphatidylethanolamine N-methyltransferase | -0.1952 | -1.1 | 0.0104341 |
| YKL006W | RPL14A | structural constituent of ribosome* | -0.1950 | -1.1 | 0.0137289 |
| YMR042W | ARG80 | DNA binding | -0.1945 | -1.1 | 0.400391 |
| YAL005C | SSA1 | ATPase * | -0.1938 | -1.1 | 0.0705759 |
| YCL011C | GBP2 | RNA binding | -0.1937 | -1.1 | 0.194575 |
| YIR032C | DAL3 | ureidoglycolate hydrolase | -0.1937 | -1.1 | 0.264595 |
| YIR013C | GAT4 | transcription factor | -0.1935 | -1.1 | 0.652147 |
| YHR039C | MSC7 | unknown | -0.1932 | -1.1 | 0.00617729 |
| YPL118W | MRP51 | structural constituent of ribosome | -0.1923 | -1.1 | 0.146772 |
| YDL060W | TSR1 | ribonucleoprotein binding | -0.1920 | -1.1 | 0.525278 |
| YJL207C | LAA1 | unknown | -0.1913 | -1.1 | 0.24327 |
| YDR110W | FOB1 | rDNA binding | -0.1912 | -1.1 | 0.519262 |
| YKL204W | EAP1 | eukaryotic initiation factor | -0.1908 | -1.1 | 0.349033 |
| YBR168W | PEX32 | unknown | -0.1907 | -1.1 | 0.0685203 |
| YNL052W | COX5A | cytochrome-c oxidase | -0.1905 | -1.1 | 0.00785718 |
| YLR241W | | unknown | -0.1903 | -1.1 | 0.162437 |
| YJL071W | ARG2 | amino-acid N-acetyltransferase | -0.1902 | -1.1 | 0.295536 |
| YDR418W | RPL12B | structural constituent of ribosome | -0.1900 | -1.1 | 0.0668711 |
| YBR218C | PYC2 | pyruvate carboxylase | -0.1898 | -1.1 | 0.0669744 |
| YOR219C | STE13 | aminopeptidase | -0.1895 | -1.1 | 0.0384164 |
| YDL113C | ATG20 | lipid binding | -0.1893 | -1.1 | 0.185725 |
| YPL005W | AEP3 | unknown | -0.1888 | -1.1 | 0.298119 |
| YPR087W | | | -0.1888 | -1.1 | 0.0253101 |
| YEL056W | HAT2 | H3/H4 histone acetyltransferase | -0.1885 | -1.1 | 0.0988706 |
| YNL082W | PMS1 | DNA binding | -0.1883 | -1.1 | 0.614609 |
| YDR342C | HXT7 | glucose transporter | -0.1883 | -1.1 | 0.0173234 |
| YJL198W | PHO90 | phosphate transporter | -0.1883 | -1.1 | 0.196866 |
| YER037W | PHM8 | unknown | -0.1882 | -1.1 | 0.159415 |
| YDR144C | MKC7 | aspartic-type signal peptidase | -0.1873 | -1.1 | 0.516103 |
| YBL005W-B | | | -0.1873 | -1.1 | 0.0699958 |
| YOL063C | CRT10 | unknown | -0.1870 | -1.1 | 0.432888 |
| YDR087C | RRP1 | unknown | -0.1868 | -1.1 | 0.192172 |
| YHR129C | ARP1 | structural constituent of cytoskeleton | -0.1867 | -1.1 | 0.14652 |
| YGR221C | TOS2 | unknown | -0.1863 | -1.1 | 0.0837577 |
| YGL167C | PMR1 | calcium-transporting ATPase * | -0.1853 | -1.1 | 0.569547 |
| YJR051W | OSM1 | fumarate reductase (NADH) | -0.1852 | -1.1 | 0.298732 |
| YOR322C | LDB19 | unknown | -0.1845 | -1.1 | 0.384461 |
| YPL048W | CAM1 | translation elongation factor | -0.1842 | -1.1 | 0.146357 |
| YDR024W | | | -0.1840 | -1.1 | 0.107408 |
| YDR057W | YOS9 | sugar binding | -0.1838 | -1.1 | 0.335856 |
| YPL174C | NIP100 | protein binding | -0.1837 | -1.1 | 0.125331 |
| YNL156C | NSG2 | unknown | -0.1832 | -1.1 | 0.0276688 |
| YFR002W | NIC96 | structural molecule | -0.1832 | -1.1 | 0.479515 |
| YDR519W | FPR2 | peptidyl-prolyl cis-trans isomerase | -0.1832 | -1.1 | 0.232466 |
| YDL128W | VCX1 | calcium ion transporter | -0.1828 | -1.1 | 0.0191666 |
| YGR210C | | unknown | -0.1828 | -1.1 | 0.430335 |
| YLL051C | FRE6 | ferric-chelate reductase | -0.1825 | -1.1 | 0.0248889 |
| YGR118W | RPS23A | structural constituent of ribosome | -0.1825 | -1.1 | 0.20151 |
| YAL049C | | unknown | -0.1823 | -1.1 | 0.120851 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YDR301W | CFT1 | RNA binding | -0.1823 | -1.1 | 0.362748 |
| YFR051C | RET2 | protein binding | -0.1822 | -1.1 | 0.614975 |
| YBR263W | SHM1 | glycine hydroxymethyltransferase | -0.1820 | -1.1 | 0.0704646 |
| YKR004C | ECM9 | unknown | -0.1808 | -1.1 | 0.187982 |
| YOL070C | | unknown | -0.1808 | -1.1 | 0.611163 |
| YHR184W | SSP1 | unknown | -0.1807 | -1.1 | 0.205008 |
| YOR080W | DIA2 | DNA replication origin binding | -0.1805 | -1.1 | 0.102328 |
| YHR207C | SET5 | unknown | -0.1805 | -1.1 | 0.175472 |
| YHR089C | GAR1 | RNA binding | -0.1802 | -1.1 | 0.254382 |
| YNL204C | SPS18 | unknown | -0.1798 | -1.1 | 0.362832 |
| YLR005W | SSL1 | general RNA polymerase II transcription factor | -0.1798 | -1.1 | 0.393944 |
| YBR217W | ATG12 | protein tag | -0.1797 | -1.1 | 0.189485 |
| YJR005W | APL1 | unknown | -0.1797 | -1.1 | 0.0916489 |
| YGL160W | | oxidoreductase | -0.1797 | -1.1 | 0.269812 |
| YBR190W | | | -0.1793 | -1.1 | 0.160566 |
| YGL078C | DBP3 | ATP-dependent RNA helicase | -0.1790 | -1.1 | 0.668114 |
| YJL202C | | | -0.1790 | -1.1 | 0.543677 |
| YJL057C | IKS1 | unknown | -0.1787 | -1.1 | 0.122885 |
| YKL146W | AVT3 | neutral amino acid transporter | -0.1787 | -1.1 | 0.466741 |
| YOR172W | YRM1 | specific RNA polymerase II transcription factor | -0.1785 | -1.1 | 0.657443 |
| YGL159W | | unknown | -0.1785 | -1.1 | 0.490604 |
| YDR505C | PSP1 | unknown | -0.1783 | -1.1 | 0.000011021 |
| YPR188C | MLC2 | myosin II binding | -0.1780 | -1.1 | 0.0613593 |
| YLR316C | TAD3 | tRNA specific adenosine deaminase | -0.1777 | -1.1 | 0.335402 |
| YDR122W | KIN1 | protein kinase | -0.1775 | -1.1 | 0.366255 |
| YIR016W | | unknown | -0.1772 | -1.1 | 0.128747 |
| YNL098C | RAS2 | GTPase | -0.1770 | -1.1 | 0.309857 |
| YNL026W | SAM50 | unknown | -0.1770 | -1.1 | 0.331834 |
| YGL144C | ROG1 | lipase | -0.1765 | -1.1 | 0.230924 |
| YFL068W | | unknown | -0.1765 | -1.1 | 0.482891 |
| YLR078C | BOS1 | v-SNARE | -0.1763 | -1.1 | 0.646854 |
| YNL209W | SSB2 | ATPase * | -0.1762 | -1.1 | 0.28421 |
| YLR069C | MEF1 | translation elongation factor | -0.1755 | -1.1 | 0.444866 |
| YDR009W | GAL3 | protein binding | -0.1752 | -1.1 | 0.401443 |
| YDR284C | DPP1 | phosphatidate phosphatase * | -0.1750 | -1.1 | 0.415261 |
| YER145C | FTR1 | iron ion transporter | -0.1750 | -1.1 | 0.683138 |
| YDR119W | | unknown | -0.1748 | -1.1 | 0.0589274 |
| YHR138C | | endopeptidase inhibitor | -0.1742 | -1.1 | 0.0384909 |
| YPR117W | | unknown | -0.1742 | -1.1 | 0.312816 |
| YEL076C | | unknown | -0.1740 | -1.1 | 0.399054 |
| YKR047W | | | -0.1735 | -1.1 | 0.207979 |
| YCRX05W | | | -0.1733 | -1.1 | 0.178291 |
| YGR242W | | | -0.1723 | -1.1 | 0.415747 |
| YER074W | RPS24A | structural constituent of ribosome | -0.1723 | -1.1 | 0.286989 |
| YKL006C-A | SFT1 | v-SNARE | -0.1723 | -1.1 | 0.207603 |
| YOL078W | AVO1 | unknown | -0.1722 | -1.1 | 0.0127619 |
| YDL124W | | alpha-keto amide reductase * | -0.1718 | -1.1 | 0.149378 |
| YER011W | TIR1 | structural constituent of cell wall | -0.1715 | -1.1 | 0.454998 |
| YML007W | YAP1 | transcription factor | -0.1715 | -1.1 | 0.0305475 |
| YLL008W | DRS1 | ATP-dependent RNA helicase | -0.1713 | -1.1 | 0.306329 |
| YHR028C | DAP2 | dipeptidyl-peptidase and tripeptidyl-peptidase | -0.1712 | -1.1 | 0.192277 |
| YNL095C | | unknown | -0.1710 | -1.1 | 0.373872 |
| YLL029W | | unknown | -0.1710 | -1.1 | 0.394528 |
| YNL241C | ZWF1 | glucose-6-phosphate 1-dehydrogenase | -0.1710 | -1.1 | 0.000651512 |
| YMR213W | CEF1 | RNA splicing factor | -0.1708 | -1.1 | 0.505728 |
| YML121W | GTR1 | GTPase | -0.1700 | -1.1 | 0.535884 |
| YJR017C | ESS1 | peptidyl-prolyl cis-trans isomerase * | -0.1695 | -1.1 | 0.0942642 |
| YEL062W | NPR2 | channel regulator | -0.1695 | -1.1 | 0.541517 |
| YML114C | TAF8 | RNA polymerase II transcription factor | -0.1690 | -1.1 | 0.563205 |
| YDR261C | EXG2 | glucan 1,3-beta-glucosidase | -0.1690 | -1.1 | 0.481691 |
| YKR094C | RPL40B | structural constituent of ribosome* | -0.1690 | -1.1 | 0.233029 |
| YER054C | GIP2 | protein phosphatase regulator | -0.1688 | -1.1 | 0.135282 |
| YPL266W | DIM1 | rRNA (adenine-N6,N6-)-dimethyltransferase | -0.1687 | -1.1 | 0.403424 |
| YOR168W | GLN4 | glutamine-tRNA ligase | -0.1685 | -1.1 | 0.0676644 |
| YBL035C | POL12 | alpha DNA polymerase | -0.1683 | -1.1 | 0.206414 |
| YCL056C | | unknown | -0.1680 | -1.1 | 0.0996936 |
| YML109W | ZDS2 | unknown | -0.1680 | -1.1 | 0.61398 |
| YKL127W | PGM1 | phosphoglucomutase | -0.1673 | -1.1 | 0.143154 |
| YIL082W-A | | | -0.1672 | -1.1 | 0.218595 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YJR155W | AAD10 | aryl-alcohol dehydrogenase | -0.1672 | -1.1 | 0.411299 |
| YKL038W | RGT1 | DNA binding | -0.1670 | -1.1 | 0.253924 |
| YDL029W | ARP2 | structural constituent of cytoskeleton* | -0.1670 | -1.1 | 0.260656 |
| YLR006C | SSK1 | enzyme activator * | -0.1668 | -1.1 | 0.0652506 |
| YGR237C | | unknown | -0.1667 | -1.1 | 0.646357 |
| YNL301C | RPL18B | structural constituent of ribosome | -0.1663 | -1.1 | 0.0604157 |
| YCR031 | | | -0.1663 | -1.1 | 0.0528392 |
| YGR055W | MUP1 | L-methionine porter | -0.1662 | -1.1 | 0.348708 |
| YFL030W | AGX1 | alanine-glyoxylate transaminase | -0.1662 | -1.1 | 0.467607 |
| YER125W | RSP5 | ubiquitin-protein ligase | -0.1660 | -1.1 | 0.011421 |
| YPL160W | CDC60 | leucine-tRNA ligase | -0.1658 | -1.1 | 0.483143 |
| YFR056C | | | -0.1658 | -1.1 | 0.143967 |
| YML079W | | unknown | -0.1655 | -1.1 | 0.357011 |
| YPL228W | CET1 | polynucleotide 5'-phosphatase | -0.1655 | -1.1 | 0.583975 |
| YGL079W | | unknown | -0.1652 | -1.1 | 0.538476 |
| YDR256C | CTA1 | catalase | -0.1652 | -1.1 | 0.0211351 |
| YOR207C | RET1 | DNA-directed RNA polymerase | -0.1648 | -1.1 | 0.436541 |
| YOL109W | ZEO1 | unknown | -0.1648 | -1.1 | 0.0374093 |
| YKR072C | SIS2 | phosphopantothienoylcysteine decarboxylase | -0.1648 | -1.1 | 0.413038 |
| YGR263C | | unknown | -0.1648 | -1.1 | 0.20384 |
| YOL026C | MIM1 | unknown | -0.1647 | -1.1 | 0.210557 |
| YLR037C | DAN2 | unknown | -0.1647 | -1.1 | 0.242491 |
| YCRX20C | | | -0.1645 | -1.1 | 0.380218 |
| YDL014W | NOP1 | methyltransferase | -0.1645 | -1.1 | 0.107702 |
| YML117W-A | | | -0.1643 | -1.1 | 0.391433 |
| YCL061C | MRC1 | unknown | -0.1643 | -1.1 | 0.21952 |
| YOR098C | NUP1 | protein binding | -0.1642 | -1.1 | 0.407653 |
| YKL097W-A | | | -0.1635 | -1.1 | 0.127224 |
| YPR103W | PRE2 | endopeptidase | -0.1632 | -1.1 | 0.303018 |
| YBR188C | NTC20 | RNA splicing factor | -0.1630 | -1.1 | 0.232823 |
| YOR326W | MYO2 | microfilament motor | -0.1630 | -1.1 | 0.573711 |
| YJR083C | ACF4 | unknown | -0.1628 | -1.1 | 0.270647 |
| YNL130C | CPT1 | diacylglycerol cholinephosphotransferase | -0.1625 | -1.1 | 0.1984 |
| YCRX04W | | | -0.1623 | -1.1 | 0.213715 |
| YDR434W | GPI17 | GPI-anchor transamidase | -0.1615 | -1.1 | 0.27259 |
| YIL097W | FYV10 | unknown | -0.1612 | -1.1 | 0.317394 |
| YDR125C | ECM18 | unknown | -0.1612 | -1.1 | 0.0848058 |
| YBR157C | ICS2 | unknown | -0.1610 | -1.1 | 0.051735 |
| YPL233W | NSL1 | unknown | -0.1607 | -1.1 | 0.516377 |
| YDR307W | | unknown | -0.1607 | -1.1 | 0.409325 |
| YGL101W | | unknown | -0.1603 | -1.1 | 0.0730395 |
| YLR097C | HRT3 | protein binding | -0.1600 | -1.1 | 0.544836 |
| YDR104C | SPO71 | unknown | -0.1600 | -1.1 | 0.0860873 |
| YOR078W | BUD21 | snoRNA binding | -0.1600 | -1.1 | 0.558052 |
| YAL059W | ECM1 | unknown | -0.1600 | -1.1 | 0.332912 |
| YLR190W | MMR1 | unknown | -0.1595 | -1.1 | 0.217316 |
| YMR099C | | unknown | -0.1590 | -1.1 | 0.162635 |
| YER077C | | unknown | -0.1588 | -1.1 | 0.0806498 |
| YNR053C | NOG2 | GTPase | -0.1588 | -1.1 | 0.678687 |
| YGL067W | NPY1 | NAD ⁺ diphosphatase | -0.1588 | -1.1 | 0.35436 |
| YER095W | RAD51 | recombinase | -0.1585 | -1.1 | 0.613685 |
| YBR115C | LYS2 | L-aminoadipate-semialdehyde dehydrogenase | -0.1583 | -1.1 | 0.213266 |
| YPL085W | SEC16 | structural molecule | -0.1582 | -1.1 | 0.724429 |
| YBR005W | RCR1 | unknown | -0.1582 | -1.1 | 0.274733 |
| YER143W | DDI1 | SNARE binding | -0.1580 | -1.1 | 0.434764 |
| YPR129W | SCD6 | RNA binding | -0.1575 | -1.1 | 0.493044 |
| YGR145W | ENP2 | unknown | -0.1573 | -1.1 | 0.375207 |
| YDR134C | | unknown | -0.1572 | -1.1 | 0.693514 |
| YOR223W | | unknown | -0.1572 | -1.1 | 0.345942 |
| YJL168C | SET2 | histone methyltransferase * | -0.1562 | -1.1 | 0.310537 |
| YLR162W | | unknown | -0.1560 | -1.1 | 0.670529 |
| YJL218W | | acetyltransferase | -0.1560 | -1.1 | 0.0773773 |
| YNL219C | ALG9 | mannosyltransferase | -0.1560 | -1.1 | 0.18093 |
| YLR429W | CRN1 | protein binding | -0.1557 | -1.1 | 0.194577 |
| YOR293W | RPS10A | structural constituent of ribosome | -0.1557 | -1.1 | 0.222946 |
| YDL050C | | | -0.1552 | -1.1 | 0.20359 |
| YLR396C | VPS33 | ATP binding | -0.1545 | -1.1 | 0.327105 |
| YOL117W | RRI2 | unknown | -0.1542 | -1.1 | 0.359962 |
| YCR053W | THR4 | threonine synthase | -0.1542 | -1.1 | 0.079656 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YBR140C | IRA1 | Ras GTPase activator | -0.1540 | -1.1 | 0.458754 |
| YDR192C | NUP42 | structural molecule | -0.1538 | -1.1 | 0.00553585 |
| YMR202W | ERG2 | C-8 sterol isomerase | -0.1532 | -1.1 | 0.171855 |
| YPL020C | ULP1 | cysteine-type peptidase * | -0.1532 | -1.1 | 0.0052059 |
| YCL019W | | | -0.1530 | -1.1 | 0.0104307 |
| YPL181W | CT16 | transcription factor | -0.1528 | -1.1 | 0.0727742 |
| YGL227W | VID30 | unknown | -0.1528 | -1.1 | 0.158456 |
| YML035C | AMD1 | AMP deaminase | -0.1525 | -1.1 | 0.200459 |
| YFL034W | | unknown | -0.1523 | -1.1 | 0.402388 |
| YDL134C-A | | | -0.1522 | -1.1 | 0.459258 |
| YKL061W | | unknown | -0.1520 | -1.1 | 0.375097 |
| YCL060C | | | -0.1518 | -1.1 | 0.466136 |
| YER019C-A | SBH2 | protein transporter | -0.1510 | -1.1 | 0.344397 |
| YDL170W | UGA3 | transcription factor | -0.1510 | -1.1 | 0.175256 |
| YPR056W | TFB4 | general RNA polymerase II transcription factor | -0.1510 | -1.1 | 0.408277 |
| YML010W | SPT5 | RNA polymerase II transcription elongation factor | -0.1510 | -1.1 | 0.150946 |
| YCR103C | | | -0.1508 | -1.1 | 0.692309 |
| YAL023C | PMT2 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.1507 | -1.1 | 0.0514421 |
| YCL028W | RNQ1 | unknown | -0.1507 | -1.1 | 0.218445 |
| YJL008C | CCT8 | unfolded protein binding | -0.1507 | -1.1 | 0.422893 |
| YCL062W | | | -0.1502 | -1.1 | 0.33935 |
| YBL039C | URA7 | CTP synthase | -0.1500 | -1.1 | 0.00835407 |
| YKL191W | DPH2 | unknown | -0.1498 | -1.1 | 0.52146 |
| YJL079C | PRY1 | unknown | -0.1495 | -1.1 | 0.108335 |
| YJL142C | | | -0.1490 | -1.1 | 0.167029 |
| YJL197W | UBP12 | ubiquitin-specific protease | -0.1490 | -1.1 | 0.174864 |
| YGR199W | PMT6 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.1487 | -1.1 | 0.658009 |
| YJR091C | JSN1 | mRNA binding | -0.1478 | -1.1 | 0.53689 |
| YCR074C | | | -0.1477 | -1.1 | 0.254702 |
| YBR176W | ECM31 | 3-methyl-2-oxobutanoate hydroxymethyltransferase | -0.1477 | -1.1 | 0.340126 |
| YOR112W | | unknown | -0.1477 | -1.1 | 0.638027 |
| YEL065W | SIT1 | siderophore-iron (ferrioxamine) uptake transporter | -0.1472 | -1.1 | 0.536128 |
| YGL105W | ARC1 | tRNA binding | -0.1472 | -1.1 | 0.419087 |
| YGR009C | SEC9 | t-SNARE | -0.1470 | -1.1 | 0.473314 |
| YJL194W | CDC6 | protein binding | -0.1470 | -1.1 | 0.506294 |
| YEL034C-A | | | -0.1468 | -1.1 | 0.0600568 |
| YJL103C | | unknown | -0.1467 | -1.1 | 0.182308 |
| YML067C | ERV41 | unknown | -0.1465 | -1.1 | 0.510663 |
| YMR264W | CUE1 | protein binding | -0.1462 | -1.1 | 0.0622459 |
| YPL013C | MRPS16 | structural constituent of ribosome | -0.1457 | -1.1 | 0.58588 |
| YAL028W | FRT2 | unknown | -0.1453 | -1.1 | 0.502288 |
| YGL198W | YIP4 | Rab GTPase binding | -0.1453 | -1.1 | 0.320778 |
| YPL244C | HUT1 | UDP-galactose transporter | -0.1452 | -1.1 | 0.719539 |
| YIL069C | RPS24B | structural constituent of ribosome | -0.1450 | -1.1 | 0.114642 |
| YNL085W | MKT1 | nuclease | -0.1447 | -1.1 | 0.783931 |
| YKR082W | NUP133 | structural molecule | -0.1445 | -1.1 | 0.58851 |
| YDL091C | UBX3 | unknown | -0.1443 | -1.1 | 0.252958 |
| YKL205W | LOS1 | tRNA binding | -0.1440 | -1.1 | 0.114825 |
| YPR058W | YMC1 | transporter | -0.1438 | -1.1 | 0.0191705 |
| YHL001W | RPL14B | structural constituent of ribosome* | -0.1432 | -1.1 | 0.0227286 |
| YBR197C | | unknown | -0.1432 | -1.1 | 0.165258 |
| YKL058W | TOA2 | general RNA polymerase II transcription factor | -0.1430 | -1.1 | 0.126217 |
| YDR463W | STP1 | transcription factor | -0.1425 | -1.1 | 0.211949 |
| YDL134C | PPH21 | protein phosphatase type 2A | -0.1423 | -1.1 | 0.239603 |
| YHR004C | NEM1 | unknown | -0.1422 | -1.1 | 0.51864 |
| YER186W-A | | | -0.1418 | -1.1 | 0.696071 |
| YDL180W | | unknown | -0.1415 | -1.1 | 0.319678 |
| YBR298C | MAL31 | alpha-glucoside:hydrogen symporter | -0.1413 | -1.1 | 0.240859 |
| YER101C | AST2 | unknown | -0.1410 | -1.1 | 0.0800624 |
| YKR005C | | unknown | -0.1410 | -1.1 | 0.743751 |
| YKR055W | RHO4 | GTPase * | -0.1408 | -1.1 | 0.120538 |
| YER063W | THO1 | unknown | -0.1407 | -1.1 | 0.280179 |
| YKL140W | TGL1 | lipase * | -0.1405 | -1.1 | 0.18071 |
| YGL162W | SUT1 | transcription factor | -0.1403 | -1.1 | 0.553877 |
| YEL015W | EDC3 | unknown | -0.1402 | -1.1 | 0.507748 |
| YDL188C | PPH22 | protein phosphatase type 2A | -0.1402 | -1.1 | 0.206989 |
| YML029W | USA1 | unknown | -0.1397 | -1.1 | 0.260129 |
| YOR084W | | lipase | -0.1397 | -1.1 | 0.4144 |
| YIL109C | SEC24 | protein binding | -0.1397 | -1.1 | 0.0170902 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YMR154C | RIM13 | cysteine-type endopeptidase | -0.1395 | -1.1 | 0.312268 |
| YPL065W | VPS28 | protein binding | -0.1393 | -1.1 | 0.244121 |
| YDL040C | NAT1 | peptide alpha-N-acetyltransferase | -0.1392 | -1.1 | 0.116916 |
| YHR056C | RSC30 | DNA binding | -0.1392 | -1.1 | 0.552839 |
| YML054C | CYB2 | L-lactate dehydrogenase | -0.1390 | -1.1 | 0.160074 |
| YER119C | AVT6 | amino acid transporter | -0.1390 | -1.1 | 0.321395 |
| YJL220W | | | -0.1387 | -1.1 | 0.502612 |
| YLR048W | RPS0B | structural constituent of ribosome | -0.1387 | -1.1 | 0.122979 |
| YKR089C | TGL4 | triacylglycerol lipase | -0.1382 | -1.1 | 0.104015 |
| YHR042W | NCP1 | electron carrier | -0.1382 | -1.1 | 0.42079 |
| YDR124W | | unknown | -0.1375 | -1.1 | 0.663955 |
| YER045C | ACA1 | specific RNA polymerase II transcription factor | -0.1373 | -1.1 | 0.517027 |
| YHR218W | | unknown | -0.1362 | -1.1 | 0.47965 |
| YKL164C | PIR1 | structural constituent of cell wall | -0.1362 | -1.1 | 0.0145756 |
| YGR206W | | unknown | -0.1355 | -1.1 | 0.25151 |
| YJL144W | | unknown | -0.1352 | -1.1 | 0.365825 |
| YPR183W | DPM1 | transferase , transferring glycosyl groups* | -0.1352 | -1.1 | 0.145994 |
| YBR224W | | | -0.1350 | -1.1 | 0.162801 |
| YDR303C | RSC3 | DNA binding | -0.1348 | -1.1 | 0.0107748 |
| YOR327C | SNC2 | v-SNARE | -0.1348 | -1.1 | 0.279553 |
| YDR210W | | unknown | -0.1347 | -1.1 | 0.027591 |
| YNL174W | | | -0.1343 | -1.1 | 0.274503 |
| YGL234W | ADE5,7 | phosphoribosylamine-glycine ligase * | -0.1343 | -1.1 | 0.614681 |
| YOL133W | HRT1 | protein binding | -0.1338 | -1.1 | 0.41228 |
| YBR052C | RFS1 | unknown | -0.1337 | -1.1 | 0.534689 |
| YHL033C | RPL8A | structural constituent of ribosome | -0.1335 | -1.1 | 0.125305 |
| YDL066W | IDP1 | isocitrate dehydrogenase | -0.1328 | -1.1 | 0.231345 |
| YER160C | | | -0.1327 | -1.1 | 0.599519 |
| YDL149W | ATG9 | unknown | -0.1327 | -1.1 | 0.507524 |
| YML082W | | carbon-sulfur lyase | -0.1325 | -1.1 | 0.367651 |
| YBR134W | | | -0.1322 | -1.1 | 0.0899691 |
| YGR168C | | unknown | -0.1322 | -1.1 | 0.561527 |
| YDR006C | SOK1 | unknown | -0.1315 | -1.1 | 0.0861411 |
| YLL022C | HIF1 | unknown | -0.1315 | -1.1 | 0.50235 |
| YGR010W | NMA2 | nicotinate-nucleotide adenyltransferase | -0.1313 | -1.1 | 0.0718076 |
| YBR081C | SPT7 | structural molecule | -0.1313 | -1.1 | 0.288386 |
| YDR012W | RPL4B | structural constituent of ribosome | -0.1312 | -1.1 | 0.18938 |
| YKR098C | UBP11 | ubiquitin-specific protease | -0.1310 | -1.1 | 0.677098 |
| YKR070W | | unknown | -0.1308 | -1.1 | 0.402678 |
| YDL042C | SIR2 | NAD-dependent histone deacetylase (H3-K14 specific)* | -0.1308 | -1.1 | 0.551522 |
| YOR032C | HMS1 | transcription factor | -0.1305 | -1.1 | 0.0376045 |
| YMR196W | | unknown | -0.1303 | -1.1 | 0.0124299 |
| YML068W | ITT1 | unknown | -0.1303 | -1.1 | 0.33264 |
| YNR051C | BRE5 | unknown | -0.1302 | -1.1 | 0.410234 |
| YBR090C | | unknown | -0.1300 | -1.1 | 0.182789 |
| YNL092W | | S-adenosylmethionine-dependent methyltransferase | -0.1297 | -1.1 | 0.525087 |
| YCRX16C | | | -0.1295 | -1.1 | 0.110964 |
| YGR248W | SOL4 | 6-phosphogluconolactonase | -0.1295 | -1.1 | 0.281456 |
| YKR003W | OSH6 | oxysterol binding | -0.1295 | -1.1 | 0.594147 |
| YER136W | GDI1 | Rab GDP-dissociation inhibitor | -0.1290 | -1.1 | 0.607331 |
| YKR031C | SPO14 | phospholipase D | -0.1288 | -1.1 | 0.362785 |
| YLR189C | ATG26 | sterol 3-beta-glucosyltransferase | -0.1288 | -1.1 | 0.130953 |
| YOL113W | SKM1 | protein serine/threonine kinase | -0.1287 | -1.1 | 0.079073 |
| YOR294W | RRS1 | unknown | -0.1282 | -1.1 | 0.156294 |
| YPL224C | MMT2 | unknown | -0.1282 | -1.1 | 0.111812 |
| YPL040C | ISM1 | isoleucine-tRNA ligase | -0.1282 | -1.1 | 0.487183 |
| YCL003W | | | -0.1282 | -1.1 | 0.310555 |
| YOR351C | MEK1 | protein serine/threonine kinase | -0.1278 | -1.1 | 0.492191 |
| YJL195C | | | -0.1273 | -1.1 | 0.429607 |
| YOR367W | SCP1 | protein binding | -0.1273 | -1.1 | 0.419922 |
| YPL190C | NAB3 | poly(A) binding | -0.1265 | -1.1 | 0.317702 |
| YGR197C | SG1 | unknown | -0.1260 | -1.1 | 0.579721 |
| YIL021W | RPB3 | DNA-directed RNA polymerase | -0.1260 | -1.1 | 0.201332 |
| YBR230C | OM14 | unknown | -0.1258 | -1.1 | 0.235694 |
| YMR131C | RRB1 | unknown | -0.1258 | -1.1 | 0.522484 |
| YFR038W | IRC5 | helicase | -0.1257 | -1.1 | 0.561434 |
| YKR087C | OMA1 | metalloendopeptidase | -0.1252 | -1.1 | 0.477111 |
| YLL043W | FPS1 | transporter | -0.1250 | -1.1 | 0.316211 |
| YJL179W | PFD1 | unfolded protein binding | -0.1248 | -1.1 | 0.234553 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|---|----------------------------|-------------|------------|
| YJR111C | | unknown | -0.1245 | -1.1 | 0.652085 |
| YHR194W | MDM31 | unknown | -0.1243 | -1.1 | 0.53552 |
| YHR168W | MTG2 | GTPase * | -0.1240 | -1.1 | 0.664784 |
| YDL164C | CDC9 | DNA ligase (ATP) | -0.1240 | -1.1 | 0.19217 |
| YIL079C | AIR1 | unknown | -0.1240 | -1.1 | 0.752631 |
| YFL005W | SEC4 | GTPase | -0.1240 | -1.1 | 0.27207 |
| YDR346C | SVF1 | unknown | -0.1237 | -1.1 | 0.379918 |
| YDR292C | SRP101 | GTP binding | -0.1237 | -1.1 | 0.166305 |
| YNR065C | YSN1 | unknown | -0.1235 | -1.1 | 0.35304 |
| YGL233W | SEC15 | protein binding | -0.1235 | -1.1 | 0.692571 |
| YNL294C | RIM21 | unknown | -0.1227 | -1.1 | 0.73693 |
| YBR041W | FAT1 | long-chain-fatty-acid-CoA ligase * | -0.1227 | -1.1 | 0.546734 |
| YDR005C | MAF1 | unknown | -0.1220 | -1.1 | 0.616001 |
| YHR073W | OSH3 | oxysterol binding | -0.1217 | -1.1 | 0.542836 |
| YHR019C | DED81 | ATP binding | -0.1217 | -1.1 | 0.558466 |
| YMR247C | RKR1 | unknown | -0.1212 | -1.1 | 0.284673 |
| YCL038C | ATG22 | unknown | -0.1212 | -1.1 | 0.383753 |
| YDR512C | EMI1 | unknown | -0.1210 | -1.1 | 0.0791264 |
| YOR142W | LSC1 | succinate-CoA ligase (ADP-forming) | -0.1208 | -1.1 | 0.250671 |
| YAL020C | ATS1 | unknown | -0.1208 | -1.1 | 0.51629 |
| YPL121C | MEI5 | unknown | -0.1207 | -1.1 | 0.432727 |
| YPL124W | SPC29 | structural constituent of cytoskeleton | -0.1205 | -1.1 | 0.120549 |
| YOL048C | | unknown | -0.1202 | -1.1 | 0.382193 |
| YPR127W | | unknown | -0.1200 | -1.1 | 0.451897 |
| YJL065C | DLS1 | unknown | -0.1198 | -1.1 | 0.483416 |
| Q0070 | AI5_ALPHA | endonuclease | -0.1197 | -1.1 | 0.456487 |
| YKL010C | UFD4 | ubiquitin-protein ligase | -0.1193 | -1.1 | 0.587727 |
| YPR115W | | unknown | -0.1192 | -1.1 | 0.663059 |
| YBR255W | | unknown | -0.1182 | -1.1 | 0.488223 |
| YDL093W | PMT5 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.1178 | -1.1 | 0.284084 |
| YPR122W | AXL1 | metalloendopeptidase | -0.1178 | -1.1 | 0.397162 |
| YIR001C | SGN1 | poly(A) binding | -0.1178 | -1.1 | 0.449523 |
| YGR279C | SCW4 | glucosidase | -0.1178 | -1.1 | 0.0277105 |
| YAR002C-A | ERP1 | unknown | -0.1173 | -1.1 | 0.701653 |
| YDR222W | | unknown | -0.1172 | -1.1 | 0.68272 |
| YGL133W | ITC1 | unknown | -0.1172 | -1.1 | 0.15743 |
| YHL035C | VMR1 | ATPase , coupled to transmembrane movement of substance | -0.1168 | -1.1 | 0.00621729 |
| YGR081C | SLX9 | unknown | -0.1167 | -1.1 | 0.634225 |
| YHL047C | ARN2 | siderophore-iron transporter | -0.1165 | -1.1 | 0.59923 |
| YDR466W | PKH3 | protein kinase | -0.1165 | -1.1 | 0.0988444 |
| YLR109W | AHP1 | thioredoxin peroxidase | -0.1162 | -1.1 | 0.129314 |
| YDR167W | TAF10 | general RNA polymerase II transcription factor | -0.1160 | -1.1 | 0.554085 |
| YLL058W | | cystathionine gamma-synthase | -0.1157 | -1.1 | 0.782135 |
| YMR068W | AVO2 | unknown | -0.1155 | -1.1 | 0.415312 |
| YJL106W | IME2 | protein kinase | -0.1153 | -1.1 | 0.664707 |
| YMR310C | | unknown | -0.1152 | -1.1 | 0.627971 |
| YBR296C | PHO89 | sodium:inorganic phosphate symporter | -0.1152 | -1.1 | 0.79518 |
| YCL061C(1) | | | -0.1152 | -1.1 | 0.401777 |
| YDR051C | | unknown | -0.1148 | -1.1 | 0.415071 |
| YIL026C | IRR1 | protein binding | -0.1145 | -1.1 | 0.288713 |
| YHR151C | | unknown | -0.1145 | -1.1 | 0.143917 |
| YKL031W | | | -0.1138 | -1.1 | 0.573415 |
| YHR130C | | | -0.1137 | -1.1 | 0.334166 |
| YGR043C | | transaldolase | -0.1135 | -1.1 | 0.419885 |
| YPL081W | RPS9A | structural constituent of ribosome | -0.1133 | -1.1 | 0.528172 |
| YJL089W | SIP4 | specific RNA polymerase II transcription factor | -0.1133 | -1.1 | 0.256606 |
| YLL003W | SFI1 | unknown | -0.1132 | -1.1 | 0.816836 |
| YNR046W | TRM112 | zinc ion binding | -0.1130 | -1.1 | 0.40181 |
| YIL142W | CCT2 | unfolded protein binding | -0.1113 | -1.1 | 0.0189217 |
| YGL021W | ALK1 | protein serine/threonine kinase | -0.1112 | -1.1 | 0.581939 |
| YGL025C | PGD1 | RNA polymerase II transcription mediator | -0.1110 | -1.1 | 0.578863 |
| YMR308C | PSE1 | protein carrier | -0.1110 | -1.1 | 0.801939 |
| YDR168W | CDC37 | unfolded protein binding | -0.1107 | -1.1 | 0.54073 |
| YKR026C | GCN3 | translation initiation factor | -0.1107 | -1.1 | 0.70457 |
| YCR023C | | unknown | -0.1103 | -1.1 | 0.503026 |
| YIR027C | DAL1 | allantoinase | -0.1100 | -1.1 | 0.35964 |
| YBR179C | FZO1 | GTPase | -0.1098 | -1.1 | 0.0903247 |
| YDR199W | | | -0.1097 | -1.1 | 0.612925 |
| YHR052W | CIC1 | protein binding | -0.1095 | -1.1 | 0.71155 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YNL318C | HXT14 | galactose transporter | -0.1090 | -1.1 | 0.561886 |
| YOR334W | MRS2 | magnesium ion transporter | -0.1090 | -1.1 | 0.475746 |
| YDL220C | CDC13 | single-stranded DNA binding | -0.1085 | -1.1 | 0.626707 |
| YLR039C | RIC1 | guanyl-nucleotide exchange factor | -0.1083 | -1.1 | 0.406453 |
| YJL213W | | unknown | -0.1083 | -1.1 | 0.0221943 |
| YOR030W | DFG16 | unknown | -0.1082 | -1.1 | 0.706998 |
| YDR132C | | unknown | -0.1080 | -1.1 | 0.396707 |
| YIL066C | RNR3 | ribonucleoside-diphosphate reductase | -0.1080 | -1.1 | 0.519733 |
| YGR149W | | unknown | -0.1078 | -1.1 | 0.58802 |
| YBR287W | ZSP1 | unknown | -0.1073 | -1.1 | 0.55854 |
| YJR075W | HOC1 | alpha-1,6-mannosyltransferase * | -0.1070 | -1.1 | 0.705919 |
| YPL219W | PCL8 | cyclin-dependent protein kinase | -0.1070 | -1.1 | 0.417464 |
| YDL115C | IWR1 | unknown | -0.1065 | -1.1 | 0.225728 |
| YIR012W | SQT1 | unknown | -0.1065 | -1.1 | 0.396582 |
| YNR031C | SSK2 | MAP kinase | -0.1065 | -1.1 | 0.641207 |
| YGR194C | XKS1 | xylulokinase | -0.1060 | -1.1 | 0.552747 |
| YPL107W | | unknown | -0.1057 | -1.1 | 0.459986 |
| YKL185W | ASH1 | specific transcriptional repressor | -0.1057 | -1.1 | 0.467262 |
| YFL007W | BLM10 | proteasome activator | -0.1055 | -1.1 | 0.640187 |
| YML102W | CAC2 | transcription regulator | -0.1053 | -1.1 | 0.385734 |
| YOR204W | DED1 | RNA helicase | -0.1053 | -1.1 | 0.394536 |
| YOR252W | TMA16 | unknown | -0.1052 | -1.1 | 0.589433 |
| YCL042W | | unknown | -0.1052 | -1.1 | 0.412876 |
| YJR086W | STE18 | GTPase | -0.1050 | -1.1 | 0.37391 |
| YCL002C | | unknown | -0.1047 | -1.1 | 0.502632 |
| YOL163W | | transporter | -0.1047 | -1.1 | 0.483146 |
| YCL068C | | unknown | -0.1047 | -1.1 | 0.417229 |
| YER076C | | unknown | -0.1043 | -1.1 | 0.545744 |
| YAL061W | | oxidoreductase | -0.1040 | -1.1 | 0.580307 |
| YFL054C | | transporter | -0.1038 | -1.1 | 0.552908 |
| YCR099C | | unknown | -0.1038 | -1.1 | 0.710152 |
| YMR222C | FSH2 | serine hydrolase | -0.1038 | -1.1 | 0.732189 |
| YJL064W | | | -0.1035 | -1.1 | 0.41045 |
| YDR499W | LCD1 | protein binding | -0.1035 | -1.1 | 0.490138 |
| YLL039C | UBI4 | protein tag* | -0.1032 | -1.1 | 0.414639 |
| YNL059C | ARP5 | unknown | -0.1028 | -1.1 | 0.540652 |
| YDR121W | DPB4 | epsilon DNA polymerase | -0.1028 | -1.1 | 0.122586 |
| YBR066C | NRG2 | transcriptional repressor | -0.1028 | -1.1 | 0.395035 |
| YBL086C | | unknown | -0.1025 | -1.1 | 0.626178 |
| Q0140 | VAR1 | structural constituent of ribosome | -0.1025 | -1.1 | 0.165831 |
| YGL166W | CUP2 | ligand-regulated transcription factor | -0.1025 | -1.1 | 0.406585 |
| YDR193W | | | -0.1023 | -1.1 | 0.562171 |
| YIL147C | SLN1 | osmosensor * | -0.1022 | -1.1 | 0.262349 |
| YCR076C | | unknown | -0.1020 | -1.1 | 0.429321 |
| YER119C-A | | | -0.1020 | -1.1 | 0.619839 |
| YGR241C | YAP1802 | clathrin binding | -0.1012 | -1.1 | 0.561593 |
| YPR081C | GRS2 | glycine-tRNA ligase | -0.1010 | -1.1 | 0.219001 |
| YDR490C | PKH1 | protein kinase | -0.1010 | -1.1 | 0.443309 |
| YDR419W | RAD30 | eta DNA polymerase | -0.1008 | -1.1 | 0.846298 |
| YFL026W | STE2 | mating-type alpha-factor | -0.1007 | -1.1 | 0.615022 |
| YJL157C | FAR1 | cyclin-dependent protein kinase | -0.1003 | -1.1 | 0.719177 |
| YCRX17W | | | -0.1002 | -1.1 | 0.566506 |
| YER036C | ARB1 | ATPase , coupled to transmembrane movement of substance | -0.1000 | -1.1 | 0.598631 |
| YOR330C | MIP1 | gamma DNA-directed DNA polymerase | -0.0998 | -1.1 | 0.737422 |
| YPL090C | RPS6A | structural constituent of ribosome | -0.0997 | -1.1 | 0.162503 |
| YEL068C | | | -0.0993 | -1.1 | 0.629579 |
| YHR197W | RIX1 | unknown | -0.0993 | -1.1 | 0.645877 |
| YDL015C | TSC13 | oxidoreductase | -0.0990 | -1.1 | 0.548753 |
| YPR021C | AGC1 | L-glutamate transporter | -0.0990 | -1.1 | 0.338194 |
| YDR478W | SNM1 | ribonuclease MRP | -0.0990 | -1.1 | 0.316946 |
| YAL048C | GEM1 | GTPase | -0.0988 | -1.1 | 0.687986 |
| YGL023C | PIB2 | phosphatidylinositol binding | -0.0988 | -1.1 | 0.446593 |
| YFR054C | | | -0.0985 | -1.1 | 0.313029 |
| YEL054C | RPL12A | structural constituent of ribosome | -0.0982 | -1.1 | 0.137389 |
| YDR460W | TFB3 | general RNA polymerase II transcription factor | -0.0978 | -1.1 | 0.208248 |
| YNR030W | ALG12 | alpha-1,6-mannosyltransferase | -0.0977 | -1.1 | 0.172108 |
| YEL002C | WBP1 | dolichyl-diphosphooligosaccharide-protein glycotransferase | -0.0977 | -1.1 | 0.481781 |
| YDR196C | | dephospho-CoA kinase | -0.0972 | -1.1 | 0.763823 |
| YAL042W | ERV46 | unknown | -0.0970 | -1.1 | 0.650228 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YJL134W | LCB3 | sphingosine-1-phosphate phosphatase | -0.0970 | -1.1 | 0.620955 |
| YPL191C | | unknown | -0.0968 | -1.1 | 0.600678 |
| YPL139C | UME1 | transcription corepressor | -0.0965 | -1.1 | 0.4321 |
| YGR214W | RPS0A | structural constituent of ribosome | -0.0960 | -1.1 | 0.41406 |
| YKR021W | | unknown | -0.0958 | -1.1 | 0.723693 |
| YDR275W | BSC2 | unknown | -0.0957 | -1.1 | 0.150502 |
| YJL080C | SCP160 | RNA binding | -0.0957 | -1.1 | 0.328908 |
| YIR014W | | unknown | -0.0953 | -1.1 | 0.655018 |
| YAL016W | TPD3 | protein phosphatase type 2A | -0.0953 | -1.1 | 0.353189 |
| YNL186W | UBP10 | ubiquitin-specific protease | -0.0953 | -1.1 | 0.330683 |
| YER017C | AFG3 | ATPase * | -0.0952 | -1.1 | 0.501967 |
| YAL021C | CCR4 | 3'-5'-exoribonuclease | -0.0950 | -1.1 | 0.460238 |
| YGR167W | CLC1 | structural molecule | -0.0948 | -1.1 | 0.600349 |
| YER108C | | | -0.0947 | -1.1 | 0.593368 |
| YNL071W | LAT1 | dihydrolipoyllysine-residue acetyltransferase | -0.0945 | -1.1 | 0.660212 |
| YHR003C | | unknown | -0.0942 | -1.1 | 0.349251 |
| YHR131C | | unknown | -0.0942 | -1.1 | 0.119275 |
| YIL155C | GUT2 | glycerol-3-phosphate dehydrogenase | -0.0942 | -1.1 | 0.552905 |
| YIL102C | | unknown | -0.0938 | -1.1 | 0.454619 |
| YDR483W | KRE2 | alpha-1,2-mannosyltransferase | -0.0933 | -1.1 | 0.157579 |
| YLR009W | RLP24 | unknown | -0.0933 | -1.1 | 0.365664 |
| YJR021C | REC107 | unknown | -0.0932 | -1.1 | 0.461811 |
| YOR141C | ARP8 | unknown | -0.0927 | -1.1 | 0.624924 |
| YML003W | | unknown | -0.0927 | -1.1 | 0.60556 |
| YBL069W | AST1 | unknown | -0.0918 | -1.1 | 0.453725 |
| YPL060C-A | | protein binding | -0.0915 | -1.1 | 0.0892925 |
| YDR240C | SNU56 | mRNA binding | -0.0915 | -1.1 | 0.740086 |
| YER079W | | unknown | -0.0912 | -1.1 | 0.666715 |
| YOL038W | PRE6 | endopeptidase | -0.0910 | -1.1 | 0.0454698 |
| YKR088C | TVP38 | unknown | -0.0910 | -1.1 | 0.378549 |
| YJL056C | ZAP1 | DNA binding | -0.0907 | -1.1 | 0.725004 |
| YDR325W | YCG1 | unknown | -0.0905 | -1.1 | 0.512063 |
| YHR132C | ECM14 | metalloendopeptidase | -0.0897 | -1.1 | 0.607188 |
| YKL124W | SSH4 | unknown | -0.0892 | -1.1 | 0.4995 |
| YGL186C | TPN1 | vitamin transporter | -0.0892 | -1.1 | 0.56691 |
| YLR206W | ENT2 | clathrin binding | -0.0888 | -1.1 | 0.566322 |
| YDL005C | MED2 | RNA polymerase II transcription mediator | -0.0882 | -1.1 | 0.210473 |
| YBR275C | RIF1 | protein binding | -0.0880 | -1.1 | 0.694292 |
| YJR006W | HYS2 | delta DNA polymerase * | -0.0878 | -1.1 | 0.466511 |
| YJL145W | SFH5 | phosphatidylinositol transporter | -0.0877 | -1.1 | 0.737913 |
| YBR227C | MCX1 | unfolded protein binding | -0.0873 | -1.1 | 0.201705 |
| YOR036W | PEP12 | t-SNARE | -0.0873 | -1.1 | 0.815163 |
| YJL158C | CIS3 | structural constituent of cell wall | -0.0870 | -1.1 | 0.552086 |
| YDR227W | SIR4 | histone binding | -0.0870 | -1.1 | 0.68597 |
| YDR388W | RVS167 | cytoskeletal protein binding | -0.0863 | -1.1 | 0.337839 |
| YBR024W | SCO2 | thioredoxin peroxidase | -0.0863 | -1.1 | 0.649241 |
| YNL214W | PEX17 | protein binding | -0.0862 | -1.1 | 0.752375 |
| YBR236C | ABD1 | mRNA (guanine-N7-)-methyltransferase | -0.0855 | -1.1 | 0.0825306 |
| YOR283W | | unknown | -0.0853 | -1.1 | 0.42211 |
| YOR335C | ALA1 | alanine-tRNA ligase | -0.0853 | -1.1 | 0.560501 |
| YKL065C | YET1 | unknown | -0.0852 | -1.1 | 0.0141304 |
| YLR025W | SNF7 | protein binding | -0.0848 | -1.1 | 0.756184 |
| YNR067C | DSE4 | glucan 1,3-beta-glucosidase | -0.0847 | -1.1 | 0.356448 |
| YER098W | UBP9 | ubiquitin-specific protease | -0.0843 | -1.1 | 0.62841 |
| YHR057C | CPR2 | peptidyl-prolyl cis-trans isomerase | -0.0842 | -1.1 | 0.387437 |
| YDL109C | | unknown | -0.0842 | -1.1 | 0.654916 |
| YHR182W | | unknown | -0.0842 | -1.1 | 0.633753 |
| YML022W | APT1 | adenine phosphoribosyltransferase | -0.0842 | -1.1 | 0.125272 |
| YJR114W | | | -0.0840 | -1.1 | 0.576313 |
| YPR088C | SRP54 | signal sequence binding | -0.0838 | -1.1 | 0.723875 |
| YLR330W | CHS5 | unknown | -0.0837 | -1.1 | 0.247817 |
| YMR076C | PDS5 | structural molecule | -0.0837 | -1.1 | 0.560107 |
| YBR158W | AMN1 | protein binding | -0.0837 | -1.1 | 0.350479 |
| YGL207W | SPT16 | RNA polymerase II transcription elongation factor | -0.0835 | -1.1 | 0.363243 |
| YJL036W | SNX4 | lipid binding | -0.0833 | -1.1 | 0.0747348 |
| YKL129C | MYO3 | microfilament motor | -0.0832 | -1.1 | 0.577019 |
| YLR433C | CNA1 | calcium-dependent protein serine/threonine phosphatase | -0.0832 | -1.1 | 0.474033 |
| YCR069W | CPR4 | peptidyl-prolyl cis-trans isomerase | -0.0832 | -1.1 | 0.407747 |
| YOR099W | KTR1 | alpha-1,2-mannosyltransferase | -0.0830 | -1.1 | 0.692675 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YPR088C | SRP54 | signal sequence binding | -0.0838 | -1.1 | 0.723875 |
| YLR330W | CHS5 | unknown | -0.0837 | -1.1 | 0.247817 |
| YMR076C | PDS5 | structural molecule | -0.0837 | -1.1 | 0.560107 |
| YBR158W | AMN1 | protein binding | -0.0837 | -1.1 | 0.350479 |
| YGL207W | SPT16 | RNA polymerase II transcription elongation factor | -0.0835 | -1.1 | 0.363243 |
| YJL036W | SNX4 | lipid binding | -0.0833 | -1.1 | 0.0747348 |
| YKL129C | MYO3 | microfilament motor | -0.0832 | -1.1 | 0.577019 |
| YLR433C | CNA1 | calcium-dependent protein serine/threonine phosphatase | -0.0832 | -1.1 | 0.474033 |
| YCR069W | CPR4 | peptidyl-prolyl cis-trans isomerase | -0.0832 | -1.1 | 0.407747 |
| YOR099W | KTR1 | alpha-1,2-mannosyltransferase | -0.0830 | -1.1 | 0.692675 |
| YIL161W | | unknown | -0.0825 | -1.1 | 0.299947 |
| YER031C | YPT31 | GTPase | -0.0825 | -1.1 | 0.524133 |
| YIL061C | SNP1 | mRNA binding | -0.0823 | -1.1 | 0.630719 |
| YIR025W | MND2 | unknown | -0.0818 | -1.1 | 0.123731 |
| YDR080W | VPS41 | Rab guanyl-nucleotide exchange factor | -0.0810 | -1.1 | 0.614169 |
| YDR120C | TRM1 | tRNA (guanine-N2-)-methyltransferase | -0.0810 | -1.1 | 0.601546 |
| YOR176W | HEM15 | ferrochelatase | -0.0807 | -1.1 | 0.0104136 |
| YHR002W | LEU5 | coenzyme A transporter | -0.0805 | -1.1 | 0.741839 |
| YMR150C | IMP1 | peptidase * | -0.0805 | -1.1 | 0.281257 |
| YPR143W | RRP15 | unknown | -0.0805 | -1.1 | 0.0625195 |
| YGL076C | RPL7A | structural constituent of ribosome | -0.0800 | -1.1 | 0.259775 |
| YGL038C | OCH1 | alpha-1,6-mannosyltransferase * | -0.0797 | -1.1 | 0.515338 |
| YAL051W | OAF1 | DNA binding | -0.0797 | -1.1 | 0.46412 |
| YKL068W | NUP100 | structural molecule | -0.0797 | -1.1 | 0.518192 |
| YNL126W | SPC98 | structural constituent of cytoskeleton | -0.0795 | -1.1 | 0.677532 |
| YBL068W | PRS4 | ribose phosphate diphosphokinase | -0.0795 | -1.1 | 0.399251 |
| YDR485C | VPS72 | unknown | -0.0795 | -1.1 | 0.490134 |
| YGL261C | PAU11 | unknown | -0.0792 | -1.1 | 0.746785 |
| YKL142W | MRP8 | structural constituent of ribosome | -0.0792 | -1.1 | 0.522486 |
| YMR295C | | unknown | -0.0792 | -1.1 | 0.0874288 |
| YNR029C | | unknown | -0.0792 | -1.1 | 0.750417 |
| YIL156W | UBP7 | ubiquitin-specific protease | -0.0790 | -1.1 | 0.471508 |
| YDR214W | AHA1 | chaperone activator | -0.0788 | -1.1 | 0.536494 |
| YDR228C | PCF11 | protein binding | -0.0788 | -1.1 | 0.415821 |
| YDR178W | SDH4 | succinate dehydrogenase | -0.0783 | -1.1 | 0.532243 |
| YJR090C | GRR1 | protein binding | -0.0782 | -1.1 | 0.301653 |
| YNL044W | YIP3 | unknown | -0.0780 | -1.1 | 0.433749 |
| YOR090C | PTC5 | protein phosphatase type 2C | -0.0780 | -1.1 | 0.418865 |
| YLR259C | HSP60 | single-stranded DNA binding | -0.0780 | -1.1 | 0.711928 |
| YNL022C | | unknown | -0.0778 | -1.1 | 0.781413 |
| YBR192W | RIM2 | transporter | -0.0777 | -1.1 | 0.383125 |
| YDR408C | ADE8 | phosphoribosylglycinamide formyltransferase | -0.0773 | -1.1 | 0.239948 |
| YMR120C | ADE17 | IMP cyclohydrolase * | -0.0770 | -1.1 | 0.545417 |
| YNL049C | SFB2 | unknown | -0.0767 | -1.1 | 0.388245 |
| YCR067C | SED4 | unknown | -0.0763 | -1.1 | 0.878061 |
| YJL033W | HCA4 | ATP-dependent RNA helicase | -0.0762 | -1.1 | 0.674849 |
| YOR350C | MNE1 | unknown | -0.0762 | -1.1 | 0.745499 |
| YGR184C | UBR1 | ubiquitin-protein ligase | -0.0755 | -1.1 | 0.586155 |
| YLL064C | PAU18 | unknown | -0.0752 | -1.1 | 0.0922036 |
| YHR115C | DMA1 | unknown | -0.0752 | -1.1 | 0.512706 |
| YNL338W | | | -0.0750 | -1.1 | 0.757544 |
| YLR461W | PAU4 | unknown | -0.0748 | -1.1 | 0.603761 |
| YKR096W | | unknown | -0.0747 | -1.1 | 0.267379 |
| YJL170C | ASG7 | unknown | -0.0742 | -1.1 | 0.685283 |
| YIL038C | NOT3 | 3'-5'-exoribonuclease | -0.0742 | -1.1 | 0.714739 |
| YIL137C | TMA108 | metalloendopeptidase | -0.0742 | -1.1 | 0.365396 |
| YLR021W | | unknown | -0.0738 | -1.1 | 0.732806 |
| YIL058W | | | -0.0732 | -1.1 | 0.706899 |
| YIR006C | PAN1 | protein binding | -0.0730 | -1.1 | 0.279951 |
| YDR276C | PMP3 | unknown | -0.0728 | -1.1 | 0.496391 |
| YGL048C | RPT6 | ATPase * | -0.0727 | -1.1 | 0.566183 |
| YGL248W | PDE1 | 3',5'-cyclic-AMP phosphodiesterase | -0.0723 | -1.1 | 0.779259 |
| YDL044C | MTF2 | RNA binding | -0.0720 | -1.1 | 0.810227 |
| YOR245C | DGA1 | diacylglycerol O-acyltransferase | -0.0718 | -1.1 | 0.436354 |
| YPL057C | SUR1 | mannosyltransferase * | -0.0712 | -1.1 | 0.592684 |
| YJR023C | | | -0.0710 | -1.1 | 0.31534 |
| YBL026W | LSM2 | RNA binding | -0.0708 | -1.1 | 0.201151 |
| YHR016C | YSC84 | unknown | -0.0708 | -1.1 | 0.704123 |
| YIL051C | MMF1 | unknown | -0.0707 | -1.1 | 0.611708 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YCR006C | | | -0.0707 | -1.1 | 0.603686 |
| YGR019W | UGA1 | 4-aminobutyrate transaminase | -0.0707 | -1.1 | 0.547897 |
| YOR355W | GDS1 | unknown | -0.0707 | -1.1 | 0.592982 |
| YMR216C | SKY1 | protein kinase | -0.0702 | -1.0 | 0.812541 |
| YKR001C | VPS1 | GTPase | -0.0698 | -1.0 | 0.790657 |
| YBR245C | ISW1 | ATPase | -0.0697 | -1.0 | 0.529826 |
| YGR218W | CRM1 | protein carrier | -0.0695 | -1.0 | 0.488509 |
| YMR285C | NGL2 | endoribonuclease | -0.0692 | -1.0 | 0.552782 |
| YLR198C | | | -0.0683 | -1.0 | 0.635873 |
| YDR142C | PEX7 | peroxisome targeting sequence binding | -0.0682 | -1.0 | 0.159493 |
| YKL032C | IXR1 | DNA binding | -0.0678 | -1.0 | 0.336766 |
| YIR035C | | unknown | -0.0677 | -1.0 | 0.794106 |
| YJR134C | SGM1 | unknown | -0.0670 | -1.0 | 0.793626 |
| YCL022C | | | -0.0670 | -1.0 | 0.74836 |
| YDR003W | RCR2 | unknown | -0.0668 | -1.0 | 0.528346 |
| YLR436C | ECM30 | unknown | -0.0667 | -1.0 | 0.764871 |
| YMR163C | | unknown | -0.0667 | -1.0 | 0.677302 |
| YBR205W | KTR3 | mannosyltransferase | -0.0667 | -1.0 | 0.465679 |
| YJR015W | | unknown | -0.0667 | -1.0 | 0.786448 |
| YIL091C | | RNA helicase | -0.0663 | -1.0 | 0.566556 |
| YJL074C | SMC3 | ATPase | -0.0658 | -1.0 | 0.72267 |
| YBR234C | ARC40 | structural constituent of cytoskeleton | -0.0658 | -1.0 | 0.329258 |
| YAL014C | SYN8 | SNAP receptor | -0.0653 | -1.0 | 0.538265 |
| YGR005C | TFG2 | general RNA polymerase II transcription factor | -0.0650 | -1.0 | 0.739137 |
| YBR209W | | | -0.0648 | -1.0 | 0.23019 |
| YML061C | PIF1 | DNA helicase | -0.0645 | -1.0 | 0.80414 |
| YCR081W | SRB8 | RNA polymerase II transcription mediator | -0.0645 | -1.0 | 0.268712 |
| YIL118W | RHO3 | GTPase * | -0.0643 | -1.0 | 0.581101 |
| YGL112C | TAF6 | general RNA polymerase II transcription factor | -0.0642 | -1.0 | 0.769054 |
| YNL302C | RPS19B | structural constituent of ribosome | -0.0642 | -1.0 | 0.212852 |
| YCL025C | AGP1 | amino acid transporter | -0.0642 | -1.0 | 0.141827 |
| YPL096W | PNG1 | peptide-N4-(N-acetyl-beta-glucosaminyl)asparagine amidas | -0.0642 | -1.0 | 0.74358 |
| YDR296W | MHR1 | transcription regulator | -0.0642 | -1.0 | 0.790316 |
| YBR260C | RGD1 | Rho GTPase activator | -0.0640 | -1.0 | 0.599352 |
| YPL138C | SPP1 | histone lysine N-methyltransferase (H3-K4 specific) | -0.0640 | -1.0 | 0.37735 |
| YLL007C | | unknown | -0.0640 | -1.0 | 0.707012 |
| YIL080W | | protein binding | -0.0638 | -1.0 | 0.213685 |
| YHR118C | ORC6 | DNA replication origin binding | -0.0638 | -1.0 | 0.759624 |
| YNL058C | | unknown | -0.0637 | -1.0 | 0.667642 |
| YOR211C | MGM1 | GTPase | -0.0637 | -1.0 | 0.756564 |
| YDR479C | PEX29 | unknown | -0.0635 | -1.0 | 0.714359 |
| YIL120W | QDR1 | multidrug transporter | -0.0633 | -1.0 | 0.419906 |
| YEL027W | CUP5 | hydrogen ion transporter | -0.0633 | -1.0 | 0.57829 |
| YDR520C | | unknown | -0.0632 | -1.0 | 0.0981678 |
| YER159C | BUR6 | transcription corepressor | -0.0632 | -1.0 | 0.589858 |
| YGR200C | ELP2 | RNA polymerase II transcription elongation factor | -0.0630 | -1.0 | 0.636785 |
| YDL077C | VAM6 | Rab guanyl-nucleotide exchange factor | -0.0628 | -1.0 | 0.642508 |
| YLR061W | RPL22A | structural constituent of ribosome | -0.0625 | -1.0 | 0.568171 |
| YJR054W | | unknown | -0.0623 | -1.0 | 0.821618 |
| YLR266C | PDR8 | DNA binding | -0.0622 | -1.0 | 0.735361 |
| YNL197C | WHI3 | RNA binding | -0.0620 | -1.0 | 0.0354348 |
| YNL230C | ELA1 | transcriptional elongation regulator | -0.0620 | -1.0 | 0.472669 |
| YPL102C | | | -0.0618 | -1.0 | 0.770348 |
| YER142C | MAG1 | alkylbase DNA N-glycosylase | -0.0617 | -1.0 | 0.663885 |
| YPR134W | MSS18 | unknown | -0.0617 | -1.0 | 0.877084 |
| YLR344W | RPL26A | structural constituent of ribosome* | -0.0615 | -1.0 | 0.529548 |
| YGL083W | SCY1 | unknown | -0.0608 | -1.0 | 0.165426 |
| YJL016W | | unknown | -0.0602 | -1.0 | 0.0949792 |
| YDR387C | | permease | -0.0598 | -1.0 | 0.736849 |
| YFR029W | PTR3 | amino acid binding | -0.0597 | -1.0 | 0.716968 |
| YIL016W | SNL1 | chaperone regulator | -0.0593 | -1.0 | 0.687521 |
| YDR106W | ARP10 | structural constituent of cytoskeleton | -0.0590 | -1.0 | 0.375597 |
| YPR101W | SNT309 | protein binding | -0.0588 | -1.0 | 0.464079 |
| YIL139C | REV7 | zeta DNA polymerase | -0.0583 | -1.0 | 0.636454 |
| YDL167C | NRP1 | unknown | -0.0583 | -1.0 | 0.688413 |
| YDL204W | RTN2 | unknown | -0.0582 | -1.0 | 0.561918 |
| YIL018W | RPL2B | structural constituent of ribosome | -0.0580 | -1.0 | 0.467008 |
| YDR271C | | | -0.0577 | -1.0 | 0.568474 |
| YBR300C | | | -0.0575 | -1.0 | 0.727731 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|----------|
| YCR055C | | | -0.0575 | -1.0 | 0.66127 |
| YDR508C | GNP1 | amino acid transporter | -0.0572 | -1.0 | 0.639641 |
| YNL330C | RPD3 | histone deacetylase | -0.0568 | -1.0 | 0.670249 |
| YIL084C | SDS3 | protein binding | -0.0563 | -1.0 | 0.571722 |
| YBL006C | LDB7 | unknown | -0.0562 | -1.0 | 0.517844 |
| YHR084W | STE12 | transcription factor | -0.0560 | -1.0 | 0.797162 |
| YPL093W | NOG1 | GTP binding | -0.0558 | -1.0 | 0.708988 |
| YOL108C | INO4 | DNA binding | -0.0557 | -1.0 | 0.807595 |
| YML110C | COQ5 | 2-hexaprenyl-6-methoxy-1,4-benzoquinone methyltransferase | -0.0557 | -1.0 | 0.438954 |
| YJR132W | NMD5 | protein carrier | -0.0555 | -1.0 | 0.550784 |
| YOL049W | GSH2 | glutathione synthase | -0.0553 | -1.0 | 0.834448 |
| YJL167W | ERG20 | dimethylallyltranstransferase * | -0.0552 | -1.0 | 0.504351 |
| YNR060W | FRE4 | ferric-chelate reductase | -0.0552 | -1.0 | 0.699472 |
| YNL035C | | unknown | -0.0550 | -1.0 | 0.901154 |
| YLR131C | ACE2 | transcriptional activator | -0.0550 | -1.0 | 0.827293 |
| YJL199C | | | -0.0550 | -1.0 | 0.33007 |
| YDR223W | CRF1 | transcriptional repressor | -0.0550 | -1.0 | 0.627539 |
| YFL035C | | | -0.0550 | -1.0 | 0.743481 |
| YLR200W | YKE2 | tubulin binding | -0.0550 | -1.0 | 0.922733 |
| YFL002C | SPB4 | ATP-dependent RNA helicase | -0.0548 | -1.0 | 0.698917 |
| YMR080C | NAM7 | ATP-dependent RNA helicase | -0.0547 | -1.0 | 0.658536 |
| YKL145W | RPT1 | ATPase * | -0.0543 | -1.0 | 0.780687 |
| YBR281C | | unknown | -0.0540 | -1.0 | 0.754943 |
| YLR362W | STE11 | MAP kinase | -0.0538 | -1.0 | 0.628554 |
| YDR194C | MSS116 | RNA splicing factor | -0.0537 | -1.0 | 0.594011 |
| YPR042C | PUF2 | mRNA binding | -0.0532 | -1.0 | 0.740833 |
| YOR010C | TIR2 | unknown | -0.0532 | -1.0 | 0.33762 |
| YPL240C | HSP82 | ATPase , coupled | -0.0528 | -1.0 | 0.656215 |
| YHL029C | OCA5 | unknown | -0.0523 | -1.0 | 0.44933 |
| YDR538W | PAD1 | carboxy-lyase | -0.0523 | -1.0 | 0.828355 |
| YIL143C | SSL2 | general RNA polymerase II transcription factor | -0.0517 | -1.0 | 0.732884 |
| YBR239C | | unknown | -0.0513 | -1.0 | 0.763219 |
| YDR062W | LCB2 | serine C-palmitoyltransferase | -0.0508 | -1.0 | 0.128936 |
| YBL024W | NCL1 | tRNA (cytosine-5-)-methyltransferase | -0.0508 | -1.0 | 0.445493 |
| YOL120C | RPL18A | structural constituent of ribosome | -0.0507 | -1.0 | 0.581717 |
| YDR415C | | unknown | -0.0507 | -1.0 | 0.408703 |
| YKL184W | SPE1 | ornithine decarboxylase | -0.0507 | -1.0 | 0.904162 |
| YPR033C | HTS1 | histidine-tRNA ligase | -0.0507 | -1.0 | 0.688431 |
| YGR250C | | RNA binding | -0.0503 | -1.0 | 0.78522 |
| YLR253W | | unknown | -0.0503 | -1.0 | 0.79846 |
| YJL099W | CHS6 | unknown | -0.0503 | -1.0 | 0.509136 |
| YAL024C | LTE1 | guanyl-nucleotide exchange factor | -0.0502 | -1.0 | 0.84551 |
| YAL031C | GIP4 | unknown | -0.0502 | -1.0 | 0.327453 |
| YDL025C | | protein kinase | -0.0500 | -1.0 | 0.715289 |
| YOR123C | LEO1 | RNA polymerase II transcription elongation factor | -0.0498 | -1.0 | 0.761871 |
| YNR038W | DBP6 | ATP-dependent RNA helicase | -0.0497 | -1.0 | 0.85949 |
| YDR052C | DBF4 | protein serine/threonine kinase | -0.0497 | -1.0 | 0.368072 |
| YJL209W | CBP1 | mRNA binding | -0.0495 | -1.0 | 0.717052 |
| YGR174C | CBP4 | unknown | -0.0495 | -1.0 | 0.494709 |
| YMR241W | YHM2 | DNA binding | -0.0493 | -1.0 | 0.822219 |
| YGR267C | FOL2 | GTP cyclohydrolase I | -0.0492 | -1.0 | 0.723074 |
| YEL038W | UTR4 | unknown | -0.0488 | -1.0 | 0.822115 |
| YBL050W | SEC17 | soluble NSF attachment protein | -0.0487 | -1.0 | 0.763212 |
| YDR055W | PST1 | unknown | -0.0487 | -1.0 | 0.645496 |
| YPR091C | | unknown | -0.0483 | -1.0 | 0.653785 |
| YOR111W | | unknown | -0.0482 | -1.0 | 0.529584 |
| YOR014W | RTS1 | protein phosphatase type 2A | -0.0478 | -1.0 | 0.634223 |
| YDR209C | | | -0.0478 | -1.0 | 0.726479 |
| YHR009C | | unknown | -0.0478 | -1.0 | 0.780936 |
| YMR045C | | | -0.0473 | -1.0 | 0.596718 |
| YKL224C | PAU16 | unknown | -0.0458 | -1.0 | 0.643853 |
| YJR065C | ARP3 | structural constituent of cytoskeleton* | -0.0458 | -1.0 | 0.805372 |
| YPL097W | MSY1 | tyrosine-tRNA ligase | -0.0458 | -1.0 | 0.856857 |
| YBR095C | RXT2 | unknown | -0.0455 | -1.0 | 0.758017 |
| YJL120W | | | -0.0452 | -1.0 | 0.744544 |
| YLR311C | | | -0.0450 | -1.0 | 0.832563 |
| YPR083W | MDM36 | unknown | -0.0448 | -1.0 | 0.880933 |
| YPL229W | | unknown | -0.0447 | -1.0 | 0.73406 |
| YLR107W | REX3 | 3'-5' exonuclease | -0.0447 | -1.0 | 0.744217 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|----------|
| YBR193C | MED8 | RNA polymerase II transcription mediator | -0.0445 | -1.0 | 0.66091 |
| YDL240W | LRG1 | Rho GTPase activator | -0.0443 | -1.0 | 0.88954 |
| YDR169C | STB3 | unknown | -0.0443 | -1.0 | 0.738794 |
| YLR027C | AAT2 | aspartate transaminase | -0.0443 | -1.0 | 0.836396 |
| YNL243W | SLA2 | protein binding | -0.0438 | -1.0 | 0.693871 |
| YLR087C | CSF1 | unknown | -0.0427 | -1.0 | 0.813665 |
| YCR077C | PAT1 | unknown | -0.0425 | -1.0 | 0.32749 |
| YPL269W | KAR9 | unknown | -0.0425 | -1.0 | 0.754015 |
| YBR279W | PAF1 | RNA polymerase II transcription elongation factor | -0.0425 | -1.0 | 0.797614 |
| YLR277C | YSH1 | RNA binding | -0.0423 | -1.0 | 0.744149 |
| YML126C | ERG13 | hydroxymethylglutaryl-CoA synthase | -0.0422 | -1.0 | 0.692601 |
| YGL030W | RPL30 | structural constituent of ribosome | -0.0418 | -1.0 | 0.77923 |
| YBL085W | BOI1 | phospholipid binding | -0.0417 | -1.0 | 0.849995 |
| YKR101W | SIR1 | DNA binding | -0.0415 | -1.0 | 0.843167 |
| YHL014C | YLF2 | GTP binding | -0.0413 | -1.0 | 0.821546 |
| YNR006W | VPS27 | protein binding | -0.0412 | -1.0 | 0.70028 |
| YDR175C | RSM24 | structural constituent of ribosome | -0.0412 | -1.0 | 0.831344 |
| YML093W | UTP14 | snoRNA binding | -0.0412 | -1.0 | 0.830397 |
| YHR165C | PRP8 | RNA splicing factor | -0.0408 | -1.0 | 0.688679 |
| YHL002W | HSE1 | protein binding | -0.0405 | -1.0 | 0.208205 |
| YDL047W | SIT4 | protein serine/threonine phosphatase | -0.0405 | -1.0 | 0.758653 |
| YOL064C | MET22 | 3'(2'),5'-biphosphate nucleotidase | -0.0398 | -1.0 | 0.899443 |
| YDR037W | KRS1 | lysine-tRNA ligase | -0.0395 | -1.0 | 0.573149 |
| YFL009W | CDC4 | protein binding | -0.0392 | -1.0 | 0.811173 |
| YLR223C | IFH1 | transcription factor | -0.0390 | -1.0 | 0.413213 |
| YGL135W | RPL1B | structural constituent of ribosome | -0.0387 | -1.0 | 0.472007 |
| YJR115W | | unknown | -0.0385 | -1.0 | 0.813272 |
| YHR032W | | unknown | -0.0385 | -1.0 | 0.881959 |
| YOR278W | HEM4 | uroporphyrinogen-III synthase | -0.0383 | -1.0 | 0.801465 |
| YHR157W | REC104 | unknown | -0.0382 | -1.0 | 0.625552 |
| YLR152C | | unknown | -0.0380 | -1.0 | 0.567681 |
| YKL108W | SLD2 | unknown | -0.0377 | -1.0 | 0.76919 |
| YCL064C | CHA1 | L-serine ammonia-lyase * | -0.0375 | -1.0 | 0.627189 |
| YDR369C | XRS2 | protein binding | -0.0373 | -1.0 | 0.771616 |
| YCR084C | TUP1 | general transcriptional repressor | -0.0373 | -1.0 | 0.692725 |
| YLR456W | | unknown | -0.0373 | -1.0 | 0.920594 |
| YAL011W | SWC3 | unknown | -0.0368 | -1.0 | 0.629277 |
| YDR309C | GIC2 | small GTPase regulator | -0.0365 | -1.0 | 0.696474 |
| YNL226W | | | -0.0365 | -1.0 | 0.910752 |
| YFR049W | YMR31 | structural constituent of ribosome | -0.0363 | -1.0 | 0.755133 |
| YIL036W | CST6 | specific RNA polymerase II transcription factor | -0.0363 | -1.0 | 0.685682 |
| YER156C | | unknown | -0.0358 | -1.0 | 0.837611 |
| YLR094C | GIS3 | unknown | -0.0357 | -1.0 | 0.428858 |
| YBL106C | SRO77 | unknown | -0.0353 | -1.0 | 0.809037 |
| YIL005W | EPS1 | protein disulfide isomerase | -0.0352 | -1.0 | 0.767351 |
| YNL199C | GCR2 | transcriptional activator | -0.0352 | -1.0 | 0.676969 |
| YNL004W | HRB1 | RNA binding | -0.0348 | -1.0 | 0.871035 |
| YLR258W | GSY2 | glycogen (starch) synthase | -0.0348 | -1.0 | 0.508692 |
| YGR192C | TDH3 | glyceraldehyde-3-phosphate dehydrogenase | -0.0348 | -1.0 | 0.906233 |
| YBR006W | UGA2 | succinate-semialdehyde dehydrogenase | -0.0348 | -1.0 | 0.827324 |
| YDR264C | AKR1 | protein-cysteine S-palmitoleyltransferase | -0.0343 | -1.0 | 0.782456 |
| YLR345W | | 6-phosphofructo-2-kinase | -0.0333 | -1.0 | 0.402839 |
| YGL259W | YPS5 | aspartic-type endopeptidase | -0.0332 | -1.0 | 0.886959 |
| YBR272C | HSM3 | unknown | -0.0328 | -1.0 | 0.801052 |
| YER038C | KRE29 | unknown | -0.0327 | -1.0 | 0.880421 |
| YFR031C | SMC2 | ATPase * | -0.0325 | -1.0 | 0.847582 |
| YJL066C | MPM1 | unknown | -0.0325 | -1.0 | 0.825287 |
| YHL017W | | unknown | -0.0318 | -1.0 | 0.738255 |
| YLR115W | CFT2 | RNA binding | -0.0315 | -1.0 | 0.864786 |
| YBL005W | PDR3 | DNA binding | -0.0315 | -1.0 | 0.877363 |
| YLR249W | YEF3 | translation elongation factor | -0.0315 | -1.0 | 0.826874 |
| YGR029W | ERV1 | thiol oxidase | -0.0315 | -1.0 | 0.910451 |
| YDR016C | DAD1 | structural constituent of cytoskeleton | -0.0312 | -1.0 | 0.711208 |
| YHR203C | RPS4B | structural constituent of ribosome | -0.0312 | -1.0 | 0.565359 |
| YML074C | FPR3 | peptidyl-prolyl cis-trans isomerase | -0.0312 | -1.0 | 0.551524 |
| YCLX11W | | | -0.0310 | -1.0 | 0.775965 |
| YMR058W | FET3 | ferroxidase | -0.0308 | -1.0 | 0.705061 |
| YDR451C | YHP1 | DNA binding | -0.0308 | -1.0 | 0.89077 |
| YDR341C | | arginine-tRNA ligase | -0.0307 | -1.0 | 0.857252 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YDL035C | GPR1 | G-protein coupled receptor | -0.0307 | -1.0 | 0.235418 |
| YEL021W | URA3 | orotidine-5'-phosphate decarboxylase | -0.0303 | -1.0 | 0.656666 |
| YJR110W | YMR1 | phosphoric monoester hydrolase * | -0.0303 | -1.0 | 0.857032 |
| YER173W | RAD24 | DNA clamp loader | -0.0302 | -1.0 | 0.819899 |
| YKL178C | STE3 | mating-type a-factor | -0.0297 | -1.0 | 0.811775 |
| YDL155W | CLB3 | cyclin-dependent protein kinase | -0.0290 | -1.0 | 0.789163 |
| YDR011W | SNQ2 | xenobiotic-transporting ATPase | -0.0283 | -1.0 | 0.512959 |
| YDR312W | SSF2 | rRNA binding | -0.0282 | -1.0 | 0.8547 |
| YDR082W | STN1 | protein binding | -0.0280 | -1.0 | 0.810832 |
| YPR036W | VMA13 | hydrogen-transporting ATPase , rotational mechanism | -0.0280 | -1.0 | 0.840109 |
| YPL272C | | unknown | -0.0278 | -1.0 | 0.902439 |
| YBR226C | | | -0.0277 | -1.0 | 0.92866 |
| YDL076C | RXT3 | unknown | -0.0270 | -1.0 | 0.629982 |
| YBR189W | RPS9B | structural constituent of ribosome | -0.0268 | -1.0 | 0.86549 |
| YPL032C | SVL3 | unknown | -0.0267 | -1.0 | 0.897005 |
| YMR125W | STO1 | mRNA binding | -0.0267 | -1.0 | 0.782313 |
| YDR170C | SEC7 | ARF guanyl-nucleotide exchange factor | -0.0267 | -1.0 | 0.680173 |
| YPL053C | KTR6 | mannosylphosphate transferase | -0.0265 | -1.0 | 0.876311 |
| YJR088C | | unknown | -0.0263 | -1.0 | 0.763243 |
| YER068W | MOT2 | 3'-5'-exoribonuclease * | -0.0262 | -1.0 | 0.909695 |
| YLR352W | | unknown | -0.0257 | -1.0 | 0.894795 |
| YBR075W | | | -0.0255 | -1.0 | 0.91607 |
| YDR471W | RPL27B | structural constituent of ribosome | -0.0255 | -1.0 | 0.806729 |
| YER005W | YND1 | nucleoside-diphosphatase | -0.0255 | -1.0 | 0.799056 |
| YLR454W | FMP27 | unknown | -0.0252 | -1.0 | 0.818101 |
| YNL314W | DAL82 | transcriptional activator | -0.0252 | -1.0 | 0.876213 |
| YDL229W | SSB1 | ATPase * | -0.0247 | -1.0 | 0.876287 |
| YDL038C | | unknown | -0.0247 | -1.0 | 0.859105 |
| YOR006C | | unknown | -0.0245 | -1.0 | 0.848252 |
| YLL027W | ISA1 | unknown | -0.0243 | -1.0 | 0.0889849 |
| YCR018C | SRD1 | unknown | -0.0240 | -1.0 | 0.882025 |
| YHR086W | NAM8 | RNA binding | -0.0238 | -1.0 | 0.905254 |
| YER018C | SPC25 | structural constituent of cytoskeleton | -0.0238 | -1.0 | 0.940266 |
| YLR002C | NOC3 | protein binding | -0.0237 | -1.0 | 0.918626 |
| YDR404C | RPB7 | DNA-directed RNA polymerase | -0.0235 | -1.0 | 0.870546 |
| YPL203W | TPK2 | protein serine/threonine kinase | -0.0233 | -1.0 | 0.849897 |
| YBR086C | IST2 | unknown | -0.0233 | -1.0 | 0.620594 |
| YER016W | BIM1 | structural constituent of cytoskeleton | -0.0232 | -1.0 | 0.895782 |
| YGL090W | LIF1 | structural molecule | -0.0230 | -1.0 | 0.873865 |
| YPL067C | | unknown | -0.0228 | -1.0 | 0.739484 |
| YLR055C | SPT8 | transcription cofactor | -0.0225 | -1.0 | 0.480433 |
| YKR038C | KAE1 | metalloendopeptidase * | -0.0222 | -1.0 | 0.884817 |
| YDL177C | | unknown | -0.0222 | -1.0 | 0.888284 |
| YGR243W | FMP43 | unknown | -0.0222 | -1.0 | 0.779583 |
| YDR211W | GCD6 | translation initiation factor | -0.0218 | -1.0 | 0.975059 |
| YFL003C | MSH4 | DNA binding | -0.0213 | -1.0 | 0.909779 |
| YHR030C | SLT2 | MAP kinase | -0.0212 | -1.0 | 0.816458 |
| YFL044C | OTU1 | ubiquitin-specific protease | -0.0212 | -1.0 | 0.739483 |
| YLR447C | VMA6 | hydrogen-transporting ATPase , rotational mechanism | -0.0207 | -1.0 | 0.84532 |
| YJL061W | NUP82 | structural molecule | -0.0205 | -1.0 | 0.699444 |
| YDL197C | ASF2 | unknown | -0.0205 | -1.0 | 0.88636 |
| YER183C | FAU1 | 5-formyltetrahydrofolate cyclo-ligase | -0.0202 | -1.0 | 0.835756 |
| YDR242W | AMD2 | amidase | -0.0202 | -1.0 | 0.79376 |
| YGL188C | | | -0.0193 | -1.0 | 0.914519 |
| YHR202W | | unknown | -0.0192 | -1.0 | 0.566001 |
| YPR004C | | electron carrier | -0.0187 | -1.0 | 0.780857 |
| YBR031W | RPL4A | structural constituent of ribosome | -0.0185 | -1.0 | 0.556407 |
| YGL215W | CLG1 | cyclin-dependent protein kinase | -0.0183 | -1.0 | 0.844932 |
| YBL092W | RPL32 | structural constituent of ribosome | -0.0183 | -1.0 | 0.901405 |
| YDR492W | IZH1 | metal ion binding | -0.0182 | -1.0 | 0.591465 |
| YIL089W | | unknown | -0.0178 | -1.0 | 0.892496 |
| YJR142W | | unknown | -0.0177 | -1.0 | 0.872164 |
| YKL116C | PRR1 | receptor signaling protein serine/threonine kinase | -0.0175 | -1.0 | 0.913189 |
| YPL070W | MUK1 | unknown | -0.0175 | -1.0 | 0.89165 |
| YIL129C | TAO3 | unknown | -0.0173 | -1.0 | 0.85467 |
| YJL160C | | unknown | -0.0170 | -1.0 | 0.851012 |
| YOR155C | ISN1 | IMP 5'-nucleotidase | -0.0165 | -1.0 | 0.89266 |
| YMR087W | | phosphoric monoester hydrolase | -0.0163 | -1.0 | 0.966298 |
| YDL119C | | transporter | -0.0163 | -1.0 | 0.725175 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|----------|
| YML028W | TSA1 | thioredoxin peroxidase | -0.0163 | -1.0 | 0.853232 |
| YNL247W | | cysteine-tRNA ligase | -0.0163 | -1.0 | 0.914835 |
| YMR182C | RGM1 | RNA polymerase II transcription factor | -0.0160 | -1.0 | 0.92509 |
| YEL043W | | unknown | -0.0160 | -1.0 | 0.933527 |
| YMR211W | DML1 | structural molecule | -0.0160 | -1.0 | 0.905066 |
| YIL123W | SIM1 | unknown | -0.0158 | -1.0 | 0.820533 |
| YDR422C | SIP1 | AMP-activated protein kinase | -0.0157 | -1.0 | 0.965271 |
| YFL019C | | | -0.0148 | -1.0 | 0.951163 |
| YOR149C | SMP3 | alpha-1,2-mannosyltransferase | -0.0147 | -1.0 | 0.750432 |
| YBL041W | PRE7 | endopeptidase | -0.0147 | -1.0 | 0.867584 |
| YHR117W | TOM71 | protein transporter | -0.0145 | -1.0 | 0.935168 |
| YLR334C | | | -0.0143 | -1.0 | 0.934978 |
| YOR362C | PRE10 | endopeptidase | -0.0142 | -1.0 | 0.881543 |
| YPL214C | THI6 | hydroxyethylthiazole kinase | -0.0142 | -1.0 | 0.882508 |
| YER099C | PRS2 | ribose phosphate diphosphokinase | -0.0140 | -1.0 | 0.833864 |
| YHR183W | GND1 | phosphogluconate dehydrogenase | -0.0140 | -1.0 | 0.885177 |
| YER118C | SHO1 | osmosensor | -0.0138 | -1.0 | 0.463847 |
| YKL029C | MAE1 | malic enzyme | -0.0138 | -1.0 | 0.63547 |
| YKL039W | PTM1 | unknown | -0.0135 | -1.0 | 0.888301 |
| YBR220C | | unknown | -0.0130 | -1.0 | 0.707317 |
| YDR076W | RAD55 | protein binding | -0.0128 | -1.0 | 0.896876 |
| YGL052W | | | -0.0128 | -1.0 | 0.961485 |
| YDR537C | | | -0.0125 | -1.0 | 0.908935 |
| YNR048W | | phospholipid-translocating ATPase | -0.0123 | -1.0 | 0.966898 |
| YBR178W | | | -0.0122 | -1.0 | 0.948954 |
| YKL094W | YJU3 | serine hydrolase | -0.0113 | -1.0 | 0.926478 |
| YLR246W | ERF2 | protein-cysteine S-palmitoleyltransferase | -0.0113 | -1.0 | 0.834724 |
| YGR007W | MUQ1 | ethanolamine-phosphate cytidyltransferase | -0.0110 | -1.0 | 0.780634 |
| YCL076W | | | -0.0108 | -1.0 | 0.937906 |
| YOL150C | | | -0.0107 | -1.0 | 0.979358 |
| YIR008C | PRI1 | alpha DNA polymerase | -0.0105 | -1.0 | 0.917847 |
| YCL010C | SGF29 | unknown | -0.0102 | -1.0 | 0.960243 |
| YBL011W | SCT1 | glycerol-3-phosphate O-acyltransferase | -0.0102 | -1.0 | 0.931716 |
| YPR016C | TIF6 | unknown | -0.0097 | -1.0 | 0.94569 |
| YLR343W | GAS2 | 1,3-beta-glucanosyltransferase | -0.0095 | -1.0 | 0.935365 |
| YMR032W | HOF1 | cytoskeletal protein binding | -0.0087 | -1.0 | 0.982998 |
| YFL021C-A | | | -0.0087 | -1.0 | 0.902364 |
| YOL009C | MDM12 | unknown | -0.0087 | -1.0 | 0.951285 |
| YBL082C | ALG3 | alpha-1,3-mannosyltransferase | -0.0085 | -1.0 | 0.961397 |
| YKL186C | MTR2 | protein binding | -0.0085 | -1.0 | 0.946552 |
| YCL040W | GLK1 | glucokinase | -0.0083 | -1.0 | 0.954952 |
| YDR108W | GSG1 | unknown | -0.0083 | -1.0 | 0.936945 |
| YPR050C | | | -0.0083 | -1.0 | 0.768144 |
| YMR100W | MUB1 | unknown | -0.0080 | -1.0 | 0.965812 |
| YMR036C | MIH1 | protein tyrosine phosphatase | -0.0080 | -1.0 | 0.969095 |
| YPL079W | RPL21B | structural constituent of ribosome | -0.0080 | -1.0 | 0.962126 |
| YOR396W | | helicase | -0.0078 | -1.0 | 0.969009 |
| YGL010W | | unknown | -0.0078 | -1.0 | 0.964464 |
| YIL067C | | unknown | -0.0067 | -1.0 | 0.987825 |
| YNL029C | KTR5 | mannosyltransferase | -0.0065 | -1.0 | 0.950516 |
| YKL141W | SDH3 | succinate dehydrogenase | -0.0065 | -1.0 | 0.972882 |
| YLR077W | FMP25 | unknown | -0.0065 | -1.0 | 0.986181 |
| YML116W | ATR1 | multidrug efflux pump | -0.0063 | -1.0 | 0.954845 |
| YHR160C | PEX18 | protein binding | -0.0063 | -1.0 | 0.968018 |
| YGL003C | CDH1 | enzyme activator | -0.0063 | -1.0 | 0.976939 |
| YAL030W | SNC1 | v-SNARE | -0.0062 | -1.0 | 0.949962 |
| YPL235W | RVB2 | ATPase * | -0.0057 | -1.0 | 0.724514 |
| YNL053W | MSG5 | prenylated protein tyrosine phosphatase | -0.0057 | -1.0 | 0.92569 |
| YJL053W | PEP8 | unknown | -0.0055 | -1.0 | 0.986169 |
| YER100W | UBC6 | ubiquitin conjugating enzyme | -0.0055 | -1.0 | 0.956651 |
| YGL228W | SHE10 | unknown | -0.0053 | -1.0 | 0.904817 |
| YEL030W | ECM10 | unknown | -0.0048 | -1.0 | 0.986114 |
| YGL194C | HOS2 | NAD-dependent histone deacetylase * | -0.0047 | -1.0 | 0.989816 |
| YOL091W | SPO21 | structural molecule | -0.0047 | -1.0 | 0.994736 |
| YLR375W | STP3 | unknown | -0.0047 | -1.0 | 0.984555 |
| YJR089W | BIR1 | unknown | -0.0045 | -1.0 | 0.989651 |
| YDR526C | | | -0.0045 | -1.0 | 0.988068 |
| YER026C | CHO1 | CDP-diacylglycerol-serine O-phosphatidyltransferase | -0.0043 | -1.0 | 0.961849 |
| YDL013W | HEX3 | DNA binding | -0.0040 | -1.0 | 0.907154 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|----------|
| YLR167W | RPS31 | structural constituent of ribosome* | -0.0038 | -1.0 | 0.953089 |
| YGL119W | ABC1 | unknown | -0.0037 | -1.0 | 0.986921 |
| YDR207C | UME6 | DNA binding | -0.0037 | -1.0 | 0.959383 |
| YOR196C | LIP5 | lipoic acid synthase | -0.0035 | -1.0 | 0.977437 |
| YNR069C | BSC5 | unknown | -0.0033 | -1.0 | 0.974219 |
| YHR140W | | unknown | -0.0030 | -1.0 | 0.974017 |
| YER089C | PTC2 | protein phosphatase type 2C | -0.0028 | -1.0 | 0.987311 |
| YEL023C | | unknown | -0.0027 | -1.0 | 0.9757 |
| YNL101W | AVT4 | neutral amino acid transporter | -0.0027 | -1.0 | 0.962791 |
| YJL084C | | cyclin binding | -0.0023 | -1.0 | 0.966938 |
| YDR259C | YAP6 | RNA polymerase II transcription factor | -0.0022 | -1.0 | 0.981948 |
| YCL008C | STP22 | protein binding | -0.0020 | -1.0 | 0.987755 |
| YKL220C | FRE2 | ferriic-chelate reductase | -0.0020 | -1.0 | 0.987206 |
| YBR181C | RPS6B | structural constituent of ribosome | -0.0018 | -1.0 | 0.985355 |
| YBR248C | HIS7 | imidazoleglycerol-phosphate synthase | -0.0017 | -1.0 | 0.98545 |
| YOR039W | CKB2 | protein kinase | -0.0015 | -1.0 | 0.9952 |
| YMR201C | RAD14 | damaged DNA binding | -0.0013 | -1.0 | 0.996328 |
| YLR148W | PEP3 | protein binding | -0.0010 | -1.0 | 0.99211 |
| YBL101C | ECM21 | unknown | -0.0010 | -1.0 | 0.977996 |
| YAL025C | MAK16 | unknown | -0.0008 | -1.0 | 0.984468 |
| YDR197W | CBS2 | unknown | -0.0003 | -1.0 | 0.997697 |
| YNL167C | SKO1 | RNA polymerase II transcription factor | -0.0003 | -1.0 | 0.996755 |
| YIL009C-A | EST3 | telomerase | -0.0002 | -1.0 | 0.998196 |
| YDR069C | DOA4 | endopeptidase * | -0.0002 | -1.0 | 0.997927 |
| YCRX19W | | | -0.0002 | -1.0 | 0.995345 |
| YFL049W | SWP82 | transcriptional activator | 0.0002 | 1.0 | 0.999866 |
| YGR038W | ORM1 | unknown | 0.0017 | 1.0 | 0.990929 |
| YHL003C | LAG1 | sphingosine N-acyltransferase | 0.0017 | 1.0 | 0.992478 |
| YJL193W | | unknown | 0.0017 | 1.0 | 0.992934 |
| YHR010W | RPL27A | structural constituent of ribosome | 0.0022 | 1.0 | 0.982863 |
| YGR282C | BGL2 | glucan 1,3-beta-glucosidase | 0.0023 | 1.0 | 0.961234 |
| YDL214C | PRR2 | receptor signaling protein serine/threonine kinase | 0.0025 | 1.0 | 0.914954 |
| YGL054C | ERV14 | unknown | 0.0027 | 1.0 | 0.983101 |
| YGL004C | RPN14 | unknown | 0.0027 | 1.0 | 0.982868 |
| YFL046W | FMP32 | unknown | 0.0027 | 1.0 | 0.976357 |
| YPL035C | | | 0.0028 | 1.0 | 0.980972 |
| YAR020C | PAU7 | unknown | 0.0032 | 1.0 | 0.977324 |
| YGR162W | TIF4631 | translation initiation factor | 0.0035 | 1.0 | 0.96052 |
| YEL076W-C | | | 0.0038 | 1.0 | 0.976404 |
| YBR124W | | | 0.0040 | 1.0 | 0.986923 |
| YBR175W | SWD3 | histone lysine N-methyltransferase (H3-K4 specific) | 0.0047 | 1.0 | 0.977356 |
| YJL129C | TRK1 | potassium ion transporter | 0.0047 | 1.0 | 0.941221 |
| YBR078W | ECM33 | unknown | 0.0047 | 1.0 | 0.801488 |
| YMR115W | FMP24 | unknown | 0.0053 | 1.0 | 0.865852 |
| YFL017C | GNA1 | glucosamine 6-phosphate N-acetyltransferase | 0.0053 | 1.0 | 0.977619 |
| YBL058W | SHP1 | protein phosphatase type 1 regulator | 0.0055 | 1.0 | 0.981329 |
| YNR033W | ABZ1 | 4-amino-4-deoxychorismate synthase | 0.0055 | 1.0 | 0.959059 |
| YHR006W | STP2 | specific RNA polymerase II transcription factor | 0.0058 | 1.0 | 0.989157 |
| YMR204C | INP1 | unknown | 0.0058 | 1.0 | 0.95237 |
| YPL131W | RPL5 | structural constituent of ribosome* | 0.0058 | 1.0 | 0.899598 |
| YIL110W | | S-adenosylmethionine-dependent methyltransferase | 0.0060 | 1.0 | 0.982015 |
| YJL100W | LSB6 | 1-phosphatidylinositol 4-kinase | 0.0063 | 1.0 | 0.971512 |
| YGL151W | NUT1 | unknown | 0.0068 | 1.0 | 0.897142 |
| YGR144W | THI4 | unknown | 0.0070 | 1.0 | 0.961283 |
| YNL151C | RPC31 | DNA-directed RNA polymerase | 0.0072 | 1.0 | 0.959531 |
| YMR325W | PAU19 | unknown | 0.0082 | 1.0 | 0.934866 |
| YCR101C | | unknown | 0.0083 | 1.0 | 0.96852 |
| YJL072C | PSF2 | unknown | 0.0083 | 1.0 | 0.964802 |
| YEL022W | GEA2 | ARF guanyl-nucleotide exchange factor | 0.0085 | 1.0 | 0.948783 |
| YOR092W | ECM3 | ATPase | 0.0090 | 1.0 | 0.971739 |
| YFR039C | | unknown | 0.0092 | 1.0 | 0.725848 |
| YGR077C | PEX8 | peroxisome targeting sequence binding | 0.0093 | 1.0 | 0.918307 |
| YLL049W | LDB18 | unknown | 0.0093 | 1.0 | 0.965144 |
| YLR289W | GUF1 | GTPase | 0.0095 | 1.0 | 0.962573 |
| YKR063C | LAS1 | unknown | 0.0097 | 1.0 | 0.809164 |
| YPL083C | SEN54 | tRNA-intron endonuclease | 0.0097 | 1.0 | 0.953372 |
| YCR019W | MAK32 | unknown | 0.0098 | 1.0 | 0.958434 |
| YKL091C | | unknown | 0.0098 | 1.0 | 0.920954 |
| YIL003W | CFD1 | ATPase | 0.0100 | 1.0 | 0.64238 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|----------|
| YPL038W | MET31 | DNA binding | 0.0100 | 1.0 | 0.947141 |
| YGR278W | CWC22 | unknown | 0.0103 | 1.0 | 0.772344 |
| YBR159W | IFA38 | ketoreductase | 0.0105 | 1.0 | 0.949024 |
| YBR229C | ROT2 | alpha-glucosidase | 0.0108 | 1.0 | 0.923785 |
| YPR054W | SMK1 | MAP kinase | 0.0112 | 1.0 | 0.935499 |
| YLR089C | ALT1 | transaminase | 0.0113 | 1.0 | 0.911161 |
| YPR065W | ROX1 | specific transcriptional repressor * | 0.0115 | 1.0 | 0.941161 |
| YML026C | RPS18B | structural constituent of ribosome | 0.0117 | 1.0 | 0.797332 |
| YDL237W | | unknown | 0.0118 | 1.0 | 0.941923 |
| YJL049W | | unknown | 0.0118 | 1.0 | 0.926511 |
| YER055C | HIS1 | ATP phosphoribosyltransferase | 0.0120 | 1.0 | 0.96253 |
| YHR054C | | unknown | 0.0122 | 1.0 | 0.866992 |
| YGL192W | IME4 | mRNA methyltransferase | 0.0123 | 1.0 | 0.960502 |
| YLR381W | CTF3 | protein binding | 0.0130 | 1.0 | 0.915219 |
| YDL142C | CRD1 | cardiolipin synthase | 0.0133 | 1.0 | 0.648914 |
| YOL132W | GAS4 | 1,3-beta-glucanosyltransferase | 0.0135 | 1.0 | 0.968147 |
| YPR055W | SEC8 | protein binding | 0.0135 | 1.0 | 0.92908 |
| YMR064W | AEP1 | unknown | 0.0143 | 1.0 | 0.407033 |
| YBR273C | UBX7 | unknown | 0.0148 | 1.0 | 0.8991 |
| YKR044W | UIP5 | unknown | 0.0150 | 1.0 | 0.860043 |
| YGL128C | CWC23 | unknown | 0.0155 | 1.0 | 0.924011 |
| YLR388W | RPS29A | structural constituent of ribosome | 0.0158 | 1.0 | 0.902194 |
| YBR051W | | | 0.0158 | 1.0 | 0.989968 |
| YNL023C | FAP1 | transcription factor | 0.0170 | 1.0 | 0.911849 |
| YNL169C | PSD1 | phosphatidylserine decarboxylase | 0.0170 | 1.0 | 0.958885 |
| YJR064W | CCT5 | unfolded protein binding | 0.0172 | 1.0 | 0.549699 |
| YLR015W | BRE2 | transcription regulator | 0.0173 | 1.0 | 0.955263 |
| YJL177W | RPL17B | structural constituent of ribosome | 0.0180 | 1.0 | 0.895936 |
| YLR410W | VIP1 | unknown | 0.0182 | 1.0 | 0.853617 |
| YLR288C | MEC3 | DNA binding | 0.0182 | 1.0 | 0.852004 |
| YHL040C | ARN1 | siderophore-iron transporter | 0.0185 | 1.0 | 0.83389 |
| YGR211W | ZPR1 | protein binding | 0.0187 | 1.0 | 0.911054 |
| YKL021C | MAK11 | unknown | 0.0188 | 1.0 | 0.829758 |
| YCRX21C | | | 0.0188 | 1.0 | 0.769287 |
| YJL159W | HSP150 | structural constituent of cell wall | 0.0192 | 1.0 | 0.541174 |
| YBR016W | | unknown | 0.0192 | 1.0 | 0.931148 |
| YML077W | BET5 | unknown | 0.0195 | 1.0 | 0.896553 |
| YIL168W | | L-serine ammonia-lyase | 0.0197 | 1.0 | 0.960346 |
| YDR277C | MTH1 | unknown | 0.0198 | 1.0 | 0.712504 |
| YKR024C | DBP7 | ATP-dependent RNA helicase | 0.0198 | 1.0 | 0.851759 |
| YPL086C | ELP3 | RNA polymerase II transcription elongation factor | 0.0203 | 1.0 | 0.700191 |
| YLR067C | PET309 | translation regulator | 0.0203 | 1.0 | 0.68054 |
| YML071C | COG8 | unknown | 0.0205 | 1.0 | 0.899939 |
| YCL016C | DCC1 | unknown | 0.0212 | 1.0 | 0.790765 |
| YPR097W | | unknown | 0.0212 | 1.0 | 0.837848 |
| YCL009C | ILV6 | enzyme regulator | 0.0213 | 1.0 | 0.434225 |
| YDL138W | RGT2 | receptor * | 0.0217 | 1.0 | 0.845873 |
| YGR288W | MAL13 | transcription factor | 0.0218 | 1.0 | 0.679269 |
| YOL032W | OPI10 | unknown | 0.0220 | 1.0 | 0.868077 |
| YHR214W | | unknown | 0.0220 | 1.0 | 0.81766 |
| YML100W | TSL1 | enzyme regulator | 0.0220 | 1.0 | 0.89818 |
| YKL166C | TPK3 | protein serine/threonine kinase | 0.0225 | 1.0 | 0.804767 |
| YLR075W | RPL10 | structural constituent of ribosome | 0.0225 | 1.0 | 0.800051 |
| YDR330W | UBX5 | unknown | 0.0228 | 1.0 | 0.908215 |
| YOL112W | MSB4 | Rab GTPase activator | 0.0230 | 1.0 | 0.895016 |
| YMR142C | RPL13B | structural constituent of ribosome | 0.0230 | 1.0 | 0.49292 |
| YKL040C | NFU1 | unknown | 0.0232 | 1.0 | 0.848228 |
| YOR238W | | unknown | 0.0233 | 1.0 | 0.937095 |
| YDL166C | FAP7 | nucleoside-triphosphatase | 0.0233 | 1.0 | 0.890104 |
| YLR347C | KAP95 | protein carrier | 0.0235 | 1.0 | 0.840648 |
| YGR217W | CCH1 | calcium channel | 0.0237 | 1.0 | 0.84024 |
| YGL081W | | unknown | 0.0238 | 1.0 | 0.775623 |
| YNL112W | DBP2 | RNA helicase | 0.0238 | 1.0 | 0.661468 |
| YLL004W | ORC3 | DNA replication origin binding | 0.0240 | 1.0 | 0.887808 |
| YDR182W | CDC1 | unknown | 0.0242 | 1.0 | 0.877942 |
| YLR066W | SPC3 | signal peptidase | 0.0247 | 1.0 | 0.230894 |
| YBR057C | MUM2 | unknown | 0.0248 | 1.0 | 0.767094 |
| YKL022C | CDC16 | protein binding | 0.0252 | 1.0 | 0.743003 |
| YOR097C | | unknown | 0.0255 | 1.0 | 0.934471 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|------------|
| YDR131C | | protein binding | 0.0260 | 1.0 | 0.915824 |
| YOR193W | PEX27 | unknown | 0.0263 | 1.0 | 0.680639 |
| YKL049C | CSE4 | centromeric DNA binding | 0.0265 | 1.0 | 0.931872 |
| YNR058W | BIO3 | adenosylmethionine-8-amino-7-oxononanoate transaminase | 0.0267 | 1.0 | 0.73044 |
| YCR072C | RSA4 | unknown | 0.0268 | 1.0 | 0.684291 |
| YGR249W | MGA1 | unknown | 0.0273 | 1.0 | 0.894769 |
| YJL081C | ARP4 | chromatin binding | 0.0275 | 1.0 | 0.784925 |
| YMR256C | COX7 | cytochrome-c oxidase | 0.0275 | 1.0 | 0.903989 |
| YNL016W | PUB1 | mRNA binding | 0.0280 | 1.0 | 0.786195 |
| YLR270W | DCS1 | hydrolase, acting on acid anhydrides, in phosphorus-contai | 0.0280 | 1.0 | 0.734285 |
| YOR125C | CAT5 | unknown | 0.0280 | 1.0 | 0.812534 |
| YMR230W | RPS10B | structural constituent of ribosome | 0.0282 | 1.0 | 0.784171 |
| YPL122C | TFB2 | general RNA polymerase II transcription factor | 0.0282 | 1.0 | 0.639985 |
| YDL033C | SLM3 | tRNA (5-methylaminomethyl-2-thiouridylate)-methyltransf | 0.0283 | 1.0 | 0.923466 |
| YBR211C | AME1 | unknown | 0.0285 | 1.0 | 0.846335 |
| YDR102C | | | 0.0285 | 1.0 | 0.767122 |
| YNR035C | ARC35 | structural molecule | 0.0287 | 1.0 | 0.755131 |
| YPR067W | ISA2 | unknown | 0.0287 | 1.0 | 0.892851 |
| YKL135C | APL2 | clathrin binding | 0.0290 | 1.0 | 0.795939 |
| YMR200W | ROT1 | unknown | 0.0292 | 1.0 | 0.760418 |
| YCR041W | | | 0.0293 | 1.0 | 0.683209 |
| YCL045C | | unknown | 0.0293 | 1.0 | 0.626402 |
| YNL176C | | unknown | 0.0293 | 1.0 | 0.867167 |
| YFL064C | | unknown | 0.0295 | 1.0 | 0.904342 |
| YDR450W | RPS18A | structural constituent of ribosome | 0.0297 | 1.0 | 0.581783 |
| YKR014C | YPT52 | GTPase | 0.0297 | 1.0 | 0.508974 |
| YDR230W | | | 0.0297 | 1.0 | 0.602258 |
| YCL065W | | | 0.0300 | 1.0 | 0.847089 |
| YDR518W | EUG1 | protein disulfide isomerase | 0.0303 | 1.0 | 0.820679 |
| YML030W | | unknown | 0.0308 | 1.0 | 0.13709 |
| YOL057W | | metalloendopeptidase | 0.0308 | 1.0 | 0.878971 |
| YEL029C | BUD16 | unknown | 0.0310 | 1.0 | 0.863763 |
| YBL057C | PTH2 | aminoacyl-tRNA hydrolase | 0.0310 | 1.0 | 0.859699 |
| YDR021W | FAL1 | ATP-dependent RNA helicase | 0.0310 | 1.0 | 0.795597 |
| YFL024C | EPL1 | histone acetyltransferase | 0.0312 | 1.0 | 0.885992 |
| YGR191W | HIP1 | L-histidine transporter | 0.0312 | 1.0 | 0.623334 |
| YDR321W | ASP1 | asparaginase | 0.0315 | 1.0 | 0.782034 |
| YJL115W | ASF1 | transcription regulator | 0.0315 | 1.0 | 0.879222 |
| YBL087C | RPL23A | structural constituent of ribosome | 0.0318 | 1.0 | 0.176756 |
| YPR005C | HAL1 | unknown | 0.0320 | 1.0 | 0.894193 |
| YDR208W | MSS4 | 1-phosphatidylinositol-4-phosphate 5-kinase | 0.0320 | 1.0 | 0.678057 |
| YBR001C | NTH2 | alpha,alpha-trehalase | 0.0323 | 1.0 | 0.838192 |
| YBR163W | DEM1 | unknown | 0.0323 | 1.0 | 0.667708 |
| YGL108C | | unknown | 0.0325 | 1.0 | 0.932601 |
| YKR037C | SPC34 | structural constituent of cytoskeleton | 0.0330 | 1.0 | 0.384864 |
| YJR014W | TMA22 | RNA binding | 0.0333 | 1.0 | 0.553566 |
| YGL223C | COG1 | unknown | 0.0337 | 1.0 | 0.914697 |
| YDR133C | | | 0.0338 | 1.0 | 0.87363 |
| YOR096W | RPS7A | structural constituent of ribosome | 0.0338 | 1.0 | 0.836781 |
| YBR235W | | ion transporter | 0.0345 | 1.0 | 0.794652 |
| YMR313C | TGL3 | triacylglycerol lipase | 0.0345 | 1.0 | 0.807244 |
| YLR421C | RPN13 | endopeptidase | 0.0348 | 1.0 | 0.723852 |
| YGR196C | FYV8 | unknown | 0.0350 | 1.0 | 0.891246 |
| YEL049W | PAU2 | unknown | 0.0352 | 1.0 | 0.474199 |
| YBL027W | RPL19B | structural constituent of ribosome | 0.0352 | 1.0 | 0.00875076 |
| YKL211C | TRP3 | anthranilate synthase * | 0.0353 | 1.0 | 0.817613 |
| YKR078W | | protein transporter | 0.0355 | 1.0 | 0.921548 |
| YDR020C | | unknown | 0.0355 | 1.0 | 0.697937 |
| YMR276W | DSK2 | protein binding | 0.0357 | 1.0 | 0.55471 |
| YGR163W | GTR2 | GTPase | 0.0357 | 1.0 | 0.874576 |
| YDR017C | KCS1 | inositol or phosphatidylinositol kinase | 0.0357 | 1.0 | 0.889864 |
| YOR220W | WSP1 | unknown | 0.0358 | 1.0 | 0.732431 |
| YOL019W | | unknown | 0.0362 | 1.0 | 0.761237 |
| YGR264C | MES1 | methionine-tRNA ligase | 0.0365 | 1.0 | 0.884875 |
| YLR392C | | unknown | 0.0365 | 1.0 | 0.83436 |
| YER048C | CAJ1 | chaperone regulator | 0.0367 | 1.0 | 0.342318 |
| YGR232W | NAS6 | unknown | 0.0368 | 1.0 | 0.858467 |
| YLR242C | ARV1 | unknown | 0.0370 | 1.0 | 0.440339 |
| YDL065C | PEX19 | protein binding | 0.0370 | 1.0 | 0.746451 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YLL031C | GPI13 | transferase , transferring phosphorus-containing groups | 0.0372 | 1.0 | 0.857006 |
| YNL138W | SRV2 | cytoskeletal protein binding | 0.0372 | 1.0 | 0.126351 |
| YDR484W | VPS52 | protein binding | 0.0377 | 1.0 | 0.791187 |
| YJL141C | YAK1 | protein kinase | 0.0380 | 1.0 | 0.325906 |
| YDR183W | PLP1 | GTPase inhibitor | 0.0382 | 1.0 | 0.838822 |
| YML112W | CTK3 | cyclin-dependent protein kinase | 0.0383 | 1.0 | 0.89062 |
| YKR020W | VPS51 | protein binding | 0.0383 | 1.0 | 0.638897 |
| YHR077C | NMD2 | protein binding | 0.0387 | 1.0 | 0.839633 |
| YER148W | SPT15 | DNA binding | 0.0387 | 1.0 | 0.839561 |
| YIL176C | PAU14 | unknown | 0.0388 | 1.0 | 0.73136 |
| YBR135W | CKS1 | protein kinase | 0.0388 | 1.0 | 0.505092 |
| YHR001W-A | QCR10 | ubiquinol-cytochrome-c reductase | 0.0392 | 1.0 | 0.769979 |
| YNR076W | PAU6 | unknown | 0.0392 | 1.0 | 0.792688 |
| YDR127W | ARO1 | 3-dehydroquinate dehydratase * | 0.0393 | 1.0 | 0.766315 |
| YBR213W | MET8 | ferrochelatase * | 0.0395 | 1.0 | 0.804685 |
| YJL091C | GWT1 | unknown | 0.0397 | 1.0 | 0.625225 |
| YAR074C | | | 0.0397 | 1.0 | 0.779955 |
| YGL210W | YPT32 | GTPase | 0.0403 | 1.0 | 0.228714 |
| YIL017W | | | 0.0405 | 1.0 | 0.652588 |
| YHR210C | | unknown | 0.0405 | 1.0 | 0.625477 |
| YKL206C | ADD66 | unknown | 0.0405 | 1.0 | 0.78834 |
| YBR288C | APM3 | unknown | 0.0408 | 1.0 | 0.638011 |
| YER056C-A | RPL34A | structural constituent of ribosome | 0.0417 | 1.0 | 0.860541 |
| YEL005C | VAB2 | protein binding | 0.0417 | 1.0 | 0.759062 |
| YPL198W | RPL7B | structural constituent of ribosome | 0.0417 | 1.0 | 0.860632 |
| YOL069W | NUF2 | structural constituent of cytoskeleton | 0.0418 | 1.0 | 0.769603 |
| YGL111W | NSA1 | unknown | 0.0420 | 1.0 | 0.847127 |
| YER042W | MXR1 | protein-methionine-S-oxide reductase | 0.0420 | 1.0 | 0.705487 |
| YLR234W | TOP3 | DNA topoisomerase type I | 0.0422 | 1.0 | 0.848958 |
| YHL036W | MUP3 | L-methionine transporter | 0.0422 | 1.0 | 0.690519 |
| YDL231C | BRE4 | unknown | 0.0422 | 1.0 | 0.721389 |
| YJL030W | MAD2 | unknown | 0.0422 | 1.0 | 0.366983 |
| YCR008W | SAT4 | protein kinase | 0.0422 | 1.0 | 0.0918535 |
| YJR016C | ILV3 | dihydroxy-acid dehydratase | 0.0423 | 1.0 | 0.461489 |
| YNL215W | IES2 | unknown | 0.0427 | 1.0 | 0.15267 |
| YBL003C | HTA2 | DNA binding | 0.0430 | 1.0 | 0.675361 |
| YKL176C | LST4 | protein transporter | 0.0430 | 1.0 | 0.070922 |
| YBR167C | POP7 | ribonuclease P * | 0.0435 | 1.0 | 0.44425 |
| YDR315C | IPK1 | inositol or phosphatidylinositol kinase | 0.0440 | 1.0 | 0.754089 |
| YKL034W | TUL1 | ubiquitin-protein ligase | 0.0440 | 1.0 | 0.58028 |
| YDL084W | SUB2 | protein binding | 0.0440 | 1.0 | 0.658628 |
| YML124C | TUB3 | structural constituent of cytoskeleton | 0.0442 | 1.0 | 0.836215 |
| YBR212W | NGR1 | RNA binding | 0.0448 | 1.0 | 0.63668 |
| YJL180C | ATP12 | unfolded protein binding | 0.0450 | 1.0 | 0.795626 |
| YDR025W | RPS11A | structural constituent of ribosome | 0.0453 | 1.0 | 0.755357 |
| YOR061W | CKA2 | protein kinase | 0.0453 | 1.0 | 0.36483 |
| YDR004W | RAD57 | protein binding | 0.0455 | 1.0 | 0.351499 |
| YPL282C | PAU22 | unknown | 0.0460 | 1.0 | 0.713519 |
| YJR145C | RPS4A | structural constituent of ribosome | 0.0462 | 1.0 | 0.707526 |
| YCR024C | SLM5 | asparagine-tRNA ligase | 0.0462 | 1.0 | 0.563609 |
| YDL216C | RR11 | metalloendopeptidase | 0.0463 | 1.0 | 0.793798 |
| YHR013C | ARD1 | peptide alpha-N-acetyltransferase | 0.0463 | 1.0 | 0.622674 |
| YGL197W | MDS3 | unknown | 0.0463 | 1.0 | 0.666317 |
| YOR132W | VPS17 | unknown | 0.0463 | 1.0 | 0.563151 |
| YOL128C | YGK3 | glycogen synthase kinase | 0.0465 | 1.0 | 0.262505 |
| YDR224C | HTB1 | DNA binding | 0.0465 | 1.0 | 0.771997 |
| YIL071W | | | 0.0472 | 1.0 | 0.44109 |
| YFL008W | SMC1 | ATPase * | 0.0472 | 1.0 | 0.136363 |
| YLR099C | ICT1 | unknown | 0.0472 | 1.0 | 0.821152 |
| YMR016C | SOK2 | transcription factor | 0.0473 | 1.0 | 0.754306 |
| YGR262C | BUD32 | protein serine/threonine kinase | 0.0475 | 1.0 | 0.800796 |
| YML106W | URA5 | orotate phosphoribosyltransferase | 0.0475 | 1.0 | 0.527079 |
| YGL035C | MIG1 | specific RNA polymerase II transcription factor | 0.0480 | 1.0 | 0.798228 |
| YCR048W | ARE1 | sterol O-acyltransferase | 0.0480 | 1.0 | 0.821126 |
| YJR151C | DAN4 | unknown | 0.0482 | 1.0 | 0.680507 |
| YPR019W | CDC54 | chromatin binding | 0.0483 | 1.0 | 0.700098 |
| YLR426W | | unknown | 0.0490 | 1.0 | 0.760096 |
| YDR148C | KGD2 | dihydrolipoyllysine-residue succinyltransferase | 0.0490 | 1.0 | 0.554161 |
| YIL013C | PDR11 | ATPase , coupled to transmembrane movement of substance | 0.0493 | 1.0 | 0.91236 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|----------|-----------|---|----------------------------|-------------|------------|
| YLL025W | PAU17 | unknown | 0.0495 | 1.0 | 0.810281 |
| YOR267C | HRK1 | protein kinase | 0.0500 | 1.0 | 0.857649 |
| YDR061W | | ATPase , coupled to transmembrane movement of substance | 0.0507 | 1.0 | 0.652089 |
| YER167W | BCK2 | unknown | 0.0508 | 1.0 | 0.710259 |
| YCL005W | LDB16 | unknown | 0.0510 | 1.0 | 0.106253 |
| YLR163C | MAS1 | mitochondrial processing peptidase | 0.0512 | 1.0 | 0.799011 |
| YPR093C | ASR1 | unknown | 0.0512 | 1.0 | 0.740813 |
| YDL058W | USO1 | unknown | 0.0515 | 1.0 | 0.72163 |
| YNL006W | LST8 | protein binding | 0.0522 | 1.0 | 0.770075 |
| YFL067W | | unknown | 0.0523 | 1.0 | 0.0332627 |
| YIL152W | | unknown | 0.0523 | 1.0 | 0.655281 |
| YER093C | TSC11 | protein binding | 0.0525 | 1.0 | 0.680729 |
| YOR143C | THI80 | thiamin diphosphokinase | 0.0527 | 1.0 | 0.778965 |
| YLR351C | NIT3 | hydrolase , acting on carbon-nitrogen (but not peptide) bonds | 0.0528 | 1.0 | 0.503414 |
| YPL022W | RAD1 | DNA binding | 0.0530 | 1.0 | 0.801398 |
| YOR148C | SPP2 | unknown | 0.0533 | 1.0 | 0.732447 |
| YBR276C | PPS1 | protein tyrosine/threonine phosphatase | 0.0533 | 1.0 | 0.321266 |
| YOL161C | PAU20 | unknown | 0.0537 | 1.0 | 0.695709 |
| YDR474C | | | 0.0543 | 1.0 | 0.659719 |
| YMR242C | RPL20A | structural constituent of ribosome | 0.0547 | 1.0 | 0.716096 |
| YMR105C | PGM2 | phosphoglucosyltransferase | 0.0548 | 1.0 | 0.580892 |
| YFR019W | FAB1 | 1-phosphatidylinositol-3-phosphate 5-kinase | 0.0550 | 1.0 | 0.501533 |
| YFR015C | GSY1 | glycogen (starch) synthase | 0.0552 | 1.0 | 0.550496 |
| YCR104W | PAU3 | unknown | 0.0557 | 1.0 | 0.424407 |
| YMR220W | ERG8 | phosphomevalonate kinase | 0.0558 | 1.0 | 0.902766 |
| YLR260W | LCB5 | D-erythro-sphingosine kinase | 0.0562 | 1.0 | 0.663162 |
| YJL078C | PRY3 | unknown | 0.0562 | 1.0 | 0.652513 |
| YDR147W | EKI1 | choline kinase | 0.0562 | 1.0 | 0.684737 |
| YJL121C | RPE1 | ribulose-phosphate 3-epimerase | 0.0562 | 1.0 | 0.361005 |
| YMR189W | GCV2 | glycine dehydrogenase | 0.0565 | 1.0 | 0.00685438 |
| YLL050C | COF1 | protein binding | 0.0567 | 1.0 | 0.404432 |
| YMR243C | ZRC1 | di-, tri-valent inorganic cation transporter | 0.0567 | 1.0 | 0.481262 |
| YPR029C | APL4 | clathrin binding | 0.0568 | 1.0 | 0.887749 |
| YLR325C | RPL38 | structural constituent of ribosome | 0.0568 | 1.0 | 0.551469 |
| YIL150C | MCM10 | chromatin binding | 0.0570 | 1.0 | 0.355852 |
| YOR320C | GNT1 | acetylglucosaminyltransferase | 0.0572 | 1.0 | 0.294937 |
| YBL019W | APN2 | phosphodiesterase I * | 0.0573 | 1.0 | 0.53149 |
| YJL140W | RPB4 | DNA-directed RNA polymerase | 0.0577 | 1.0 | 0.493736 |
| YOR043W | WHI2 | phosphatase activator | 0.0578 | 1.0 | 0.525622 |
| YMR194W | RPL36A | structural constituent of ribosome* | 0.0580 | 1.0 | 0.238934 |
| YOR250C | CLP1 | RNA binding | 0.0580 | 1.0 | 0.884374 |
| YMR088C | VBA1 | basic amino acid permease | 0.0583 | 1.0 | 0.849485 |
| YOR246C | | oxidoreductase | 0.0583 | 1.0 | 0.118339 |
| YOR332W | VMA4 | hydrogen-transporting ATPase , rotational mechanism | 0.0583 | 1.0 | 0.617465 |
| YJL014W | CCT3 | unfolded protein binding | 0.0587 | 1.0 | 0.455095 |
| YNL266W | | | 0.0590 | 1.0 | 0.6418 |
| YDR068W | DOS2 | unknown | 0.0592 | 1.0 | 0.64928 |
| YBR161W | CSH1 | mannosyltransferase * | 0.0593 | 1.0 | 0.684661 |
| YMR261C | TPS3 | enzyme regulator | 0.0593 | 1.0 | 0.0444578 |
| YPL023C | MET12 | methylenetetrahydrofolate reductase (NADPH) | 0.0593 | 1.0 | 0.807659 |
| YOL027C | MDM38 | unknown | 0.0598 | 1.0 | 0.453741 |
| YBR133C | HSL7 | protein-arginine N-methyltransferase | 0.0598 | 1.0 | 0.135992 |
| YJL013C | MAD3 | unknown | 0.0600 | 1.0 | 0.483097 |
| YDR162C | NBP2 | unknown | 0.0605 | 1.0 | 0.706345 |
| YFL065C | | unknown | 0.0610 | 1.0 | 0.694884 |
| YPR102C | RPL11A | structural constituent of ribosome | 0.0613 | 1.0 | 0.0491071 |
| YNL119W | NCS2 | unknown | 0.0617 | 1.0 | 0.739047 |
| YIL095W | PRK1 | protein serine/threonine kinase | 0.0620 | 1.0 | 0.520303 |
| YOR359W | VTS1 | RNA binding | 0.0620 | 1.0 | 0.272477 |
| YMR218C | TRS130 | unknown | 0.0625 | 1.0 | 0.647464 |
| YHR188C | GPI16 | GPI-anchor transamidase | 0.0633 | 1.0 | 0.642159 |
| YIR041W | PAU15 | unknown | 0.0633 | 1.0 | 0.624994 |
| YDR109C | | kinase | 0.0635 | 1.0 | 0.492814 |
| YPL149W | ATG5 | unknown | 0.0635 | 1.0 | 0.676752 |
| YLR342W | FKS1 | 1,3-beta-glucan synthase | 0.0638 | 1.0 | 0.782619 |
| YGL137W | SEC27 | unknown | 0.0638 | 1.0 | 0.764248 |
| YDR105C | TMS1 | unknown | 0.0640 | 1.0 | 0.268727 |
| YCL030C | HIS4 | phosphoribosyl-ATP diphosphatase * | 0.0640 | 1.0 | 0.583129 |
| YCR097WA | | | 0.0642 | 1.0 | 0.685302 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YKL046C | DCW1 | mannosidase | 0.0643 | 1.0 | 0.754956 |
| YJL035C | TAD2 | tRNA specific adenosine deaminase | 0.0643 | 1.0 | 0.376867 |
| YJL139C | YUR1 | mannosyltransferase | 0.0647 | 1.0 | 0.551196 |
| YJL127C | SPT10 | histone acetyltransferase | 0.0648 | 1.0 | 0.286744 |
| YOR383C | FIT3 | unknown | 0.0650 | 1.0 | 0.579344 |
| YKL047W | | unknown | 0.0652 | 1.0 | 0.83548 |
| YGL085W | | unknown | 0.0653 | 1.0 | 0.46591 |
| YBR112C | CYC8 | transcription coactivator * | 0.0655 | 1.0 | 0.122959 |
| YDL010W | | unknown | 0.0658 | 1.0 | 0.499612 |
| YOR144C | ELG1 | unknown | 0.0658 | 1.0 | 0.851304 |
| YOR089C | VPS21 | GTPase | 0.0660 | 1.0 | 0.543415 |
| YER181C | | | 0.0667 | 1.0 | 0.594153 |
| YKL122C | SRP21 | signal sequence binding | 0.0668 | 1.0 | 0.447795 |
| YOR103C | OST2 | dolichyl-diphosphooligosaccharide-protein glycotransferase | 0.0670 | 1.0 | 0.690893 |
| YGR102C | | unknown | 0.0672 | 1.0 | 0.197037 |
| YPL044C | | | 0.0673 | 1.0 | 0.799545 |
| YER082C | UTP7 | snoRNA binding | 0.0675 | 1.0 | 0.450726 |
| YNL259C | ATX1 | copper chaperone | 0.0675 | 1.0 | 0.671987 |
| YPR079W | MRL1 | signal sequence binding | 0.0675 | 1.0 | 0.827861 |
| YDR091C | RLI1 | ATPase * | 0.0677 | 1.0 | 0.560033 |
| YNL225C | CNM67 | structural constituent of cytoskeleton | 0.0677 | 1.0 | 0.718632 |
| YER109C | FLO8 | specific RNA polymerase II transcription factor | 0.0678 | 1.0 | 0.483187 |
| YNL254C | | unknown | 0.0678 | 1.0 | 0.699391 |
| YMR255W | GFD1 | unknown | 0.0680 | 1.0 | 0.485423 |
| YML073C | RPL6A | structural constituent of ribosome* | 0.0680 | 1.0 | 0.361732 |
| YBR102C | EXO84 | protein binding | 0.0682 | 1.0 | 0.495793 |
| YBR058C | UBP14 | ubiquitin-specific protease | 0.0682 | 1.0 | 0.782692 |
| YBL102W | SFT2 | unknown | 0.0687 | 1.0 | 0.611863 |
| YER186C | | unknown | 0.0688 | 1.0 | 0.672433 |
| YIL017C | VID28 | unknown | 0.0688 | 1.0 | 0.813693 |
| YGR070W | ROM1 | signal transducer * | 0.0693 | 1.0 | 0.764803 |
| YGL092W | NUP145 | structural molecule | 0.0695 | 1.0 | 0.831691 |
| YHL002C-A | | | 0.0697 | 1.0 | 0.00904149 |
| YDR372C | VPS74 | unknown | 0.0700 | 1.0 | 0.13573 |
| YKL048C | ELM1 | protein serine/threonine kinase | 0.0708 | 1.1 | 0.412275 |
| YPL273W | SAM4 | homocysteine S-methyltransferase | 0.0710 | 1.1 | 0.586741 |
| YER004W | FMP52 | unknown | 0.0710 | 1.1 | 0.653063 |
| YDR263C | DIN7 | nuclease | 0.0713 | 1.1 | 0.22303 |
| YDL176W | | unknown | 0.0713 | 1.1 | 0.696016 |
| YLR144C | ACF2 | glucan 1,3-beta-glucosidase | 0.0717 | 1.1 | 0.748828 |
| YHL042W | | unknown | 0.0717 | 1.1 | 0.39104 |
| YPL071C | | unknown | 0.0718 | 1.1 | 0.286152 |
| YDR516C | EMI2 | unknown | 0.0725 | 1.1 | 0.571553 |
| YGR132C | PHB1 | unknown | 0.0732 | 1.1 | 0.325858 |
| YDR428C | | unknown | 0.0732 | 1.1 | 0.657755 |
| YLR233C | EST1 | RNA binding | 0.0735 | 1.1 | 0.810466 |
| YKL051W | SFK1 | unknown | 0.0738 | 1.1 | 0.214504 |
| YBR079C | RPG1 | translation initiation factor | 0.0742 | 1.1 | 0.283855 |
| YAL019W | FUN30 | unknown | 0.0742 | 1.1 | 0.69321 |
| YDL112W | TRM3 | tRNA (guanosine) methyltransferase | 0.0747 | 1.1 | 0.29187 |
| YIL134W | FLX1 | FAD transporter | 0.0748 | 1.1 | 0.67789 |
| YLR301W | | unknown | 0.0752 | 1.1 | 0.656596 |
| YOR027W | STI1 | unfolded protein binding | 0.0755 | 1.1 | 0.549152 |
| YDR060W | MAK21 | unknown | 0.0755 | 1.1 | 0.117159 |
| YLR312C | | unknown | 0.0755 | 1.1 | 0.437697 |
| YDL063C | | unknown | 0.0758 | 1.1 | 0.318933 |
| YBR231C | SWC5 | unknown | 0.0762 | 1.1 | 0.475621 |
| YER046W | SPO73 | unknown | 0.0763 | 1.1 | 0.445665 |
| YBR017C | KAP104 | nuclear localization sequence binding | 0.0763 | 1.1 | 0.651697 |
| YPL217C | BMS1 | GTP binding | 0.0765 | 1.1 | 0.586416 |
| YGL063W | PUS2 | tRNA-pseudouridine synthase | 0.0767 | 1.1 | 0.693064 |
| YPL234C | TFP3 | hydrogen-transporting ATPase , rotational mechanism | 0.0767 | 1.1 | 0.524751 |
| YIL148W | RPL40A | structural constituent of ribosome* | 0.0770 | 1.1 | 0.1342 |
| YLR177W | | unknown | 0.0770 | 1.1 | 0.772262 |
| YDL064W | UBC9 | SUMO conjugating enzyme | 0.0773 | 1.1 | 0.20966 |
| YPL179W | PPQ1 | protein serine/threonine phosphatase | 0.0775 | 1.1 | 0.435543 |
| YMR300C | ADE4 | amidophosphoribosyltransferase | 0.0777 | 1.1 | 0.601431 |
| YBR247C | ENP1 | snoRNA binding | 0.0778 | 1.1 | 0.452546 |
| YCR100C | | unknown | 0.0778 | 1.1 | 0.647394 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YGR112W | SHY1 | unfolded protein binding | 0.0782 | 1.1 | 0.338026 |
| YMR033W | ARP9 | general RNA polymerase II transcription factor | 0.0782 | 1.1 | 0.737167 |
| YGR049W | SCM4 | unknown | 0.0785 | 1.1 | 0.436405 |
| YLR215C | CDC123 | unknown | 0.0785 | 1.1 | 0.646919 |
| YKR062W | TFA2 | general RNA polymerase II transcription factor | 0.0785 | 1.1 | 0.0699522 |
| YNL178W | RPS3 | structural constituent of ribosome | 0.0785 | 1.1 | 0.499495 |
| YMR289W | ABZ2 | unknown | 0.0787 | 1.1 | 0.632771 |
| YLR047C | FRE8 | oxidoreductase | 0.0790 | 1.1 | 0.840524 |
| YGR186W | TFG1 | general RNA polymerase II transcription factor | 0.0795 | 1.1 | 0.801059 |
| YMR077C | VPS20 | protein binding | 0.0797 | 1.1 | 0.0991138 |
| YKR071C | DRE2 | unknown | 0.0798 | 1.1 | 0.414462 |
| YCR015C | | unknown | 0.0800 | 1.1 | 0.502485 |
| YIL048W | NEO1 | phospholipid-translocating ATPase * | 0.0807 | 1.1 | 0.346581 |
| YMR074C | | unknown | 0.0810 | 1.1 | 0.54994 |
| YDR294C | DPL1 | sphinganine-1-phosphate aldolase | 0.0815 | 1.1 | 0.526139 |
| YBR237W | PRP5 | RNA splicing factor | 0.0815 | 1.1 | 0.236781 |
| YLR408C | | unknown | 0.0818 | 1.1 | 0.731278 |
| YPR043W | RPL43A | structural constituent of ribosome | 0.0822 | 1.1 | 0.195287 |
| YDL196W | | | 0.0822 | 1.1 | 0.472237 |
| YGL181W | GTS1 | unknown | 0.0823 | 1.1 | 0.420337 |
| YHL049C | | unknown | 0.0827 | 1.1 | 0.454649 |
| YOR312C | RPL20B | structural constituent of ribosome | 0.0827 | 1.1 | 0.0348962 |
| YHR078W | | unknown | 0.0828 | 1.1 | 0.182504 |
| YHR011W | DIA4 | serine-tRNA ligase | 0.0833 | 1.1 | 0.0705088 |
| YKL222C | | unknown | 0.0835 | 1.1 | 0.693354 |
| YJR013W | GPI14 | mannosyltransferase | 0.0837 | 1.1 | 0.607683 |
| YGL009C | LEU1 | 3-isopropylmalate dehydratase | 0.0838 | 1.1 | 0.79821 |
| YDR185C | | unknown | 0.0845 | 1.1 | 0.353996 |
| YCL073C | | transporter | 0.0848 | 1.1 | 0.549888 |
| YMR060C | SAM37 | protein binding | 0.0848 | 1.1 | 0.669086 |
| YOR117W | RPT5 | ATPase * | 0.0850 | 1.1 | 0.690479 |
| YBR238C | | unknown | 0.0852 | 1.1 | 0.371894 |
| YPL144W | | unknown | 0.0852 | 1.1 | 0.354544 |
| YHR201C | PPX1 | exopolyphosphatase | 0.0855 | 1.1 | 0.322964 |
| YKL207W | | unknown | 0.0857 | 1.1 | 0.550222 |
| YDR101C | ARX1 | unknown | 0.0865 | 1.1 | 0.597683 |
| YIL011W | TIR3 | unknown | 0.0867 | 1.1 | 0.161438 |
| YER049W | TPA1 | unknown | 0.0868 | 1.1 | 0.623126 |
| YDL141W | BPL1 | biotin-[acetyl-CoA-carboxylase] ligase * | 0.0870 | 1.1 | 0.252365 |
| YOR272W | YTM1 | unknown | 0.0870 | 1.1 | 0.252762 |
| YKL162C | | unknown | 0.0872 | 1.1 | 0.367748 |
| YMR069W | NAT4 | peptide alpha-N-acetyltransferase | 0.0872 | 1.1 | 0.679046 |
| YBR216C | YBP1 | unknown | 0.0872 | 1.1 | 0.478932 |
| YGR258C | RAD2 | single-stranded DNA specific endodeoxyribonuclease | 0.0873 | 1.1 | 0.579613 |
| YCL018W | LEU2 | 3-isopropylmalate dehydrogenase | 0.0878 | 1.1 | 0.0434134 |
| YBR110W | ALG1 | beta-1,4-mannosyltransferase | 0.0880 | 1.1 | 0.360102 |
| YPL095C | EEB1 | unknown | 0.0882 | 1.1 | 0.506272 |
| YOL039W | RPP2A | structural constituent of ribosome | 0.0887 | 1.1 | 0.181571 |
| YER184C | | unknown | 0.0888 | 1.1 | 0.661686 |
| YDL224C | WHI4 | RNA binding | 0.0888 | 1.1 | 0.445724 |
| YDR293C | SSD1 | RNA binding | 0.0892 | 1.1 | 0.428122 |
| YIL022W | TIM44 | protein transporter | 0.0892 | 1.1 | 0.70065 |
| YDR487C | RIB3 | 3,4-dihydroxy-2-butanone-4-phosphate synthase | 0.0893 | 1.1 | 0.0209302 |
| YER179W | DMC1 | single-stranded DNA binding | 0.0893 | 1.1 | 0.626534 |
| YDL189W | RBS1 | unknown | 0.0895 | 1.1 | 0.11124 |
| YIL135C | VHS2 | unknown | 0.0895 | 1.1 | 0.352168 |
| YJR079W | | unknown | 0.0897 | 1.1 | 0.70391 |
| YDR019C | GCV1 | glycine dehydrogenase | 0.0898 | 1.1 | 0.77012 |
| YDR137W | RGP1 | guanyl-nucleotide exchange factor | 0.0898 | 1.1 | 0.667073 |
| YPL098C | MGR2 | unknown | 0.0900 | 1.1 | 0.535491 |
| YML051W | GAL80 | transcription corepressor | 0.0903 | 1.1 | 0.191366 |
| YGL178W | MPT5 | mRNA binding | 0.0905 | 1.1 | 0.769038 |
| YDL088C | ASM4 | structural molecule | 0.0907 | 1.1 | 0.488145 |
| YBR044C | TCM62 | unfolded protein binding | 0.0908 | 1.1 | 0.606928 |
| YDR258C | HSP78 | ATPase * | 0.0910 | 1.1 | 0.0854136 |
| YCL032W | STE50 | protein kinase | 0.0913 | 1.1 | 0.576846 |
| YGR111W | | unknown | 0.0915 | 1.1 | 0.254635 |
| YNL127W | FAR11 | unknown | 0.0915 | 1.1 | 0.484484 |
| YGL172W | NUP49 | structural molecule | 0.0920 | 1.1 | 0.676493 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YER010C | | unknown | 0.0922 | 1.1 | 0.579126 |
| YOR270C | VPH1 | hydrogen-transporting ATPase , rotational mechanism | 0.0922 | 1.1 | 0.574421 |
| YFL033C | RIM15 | protein kinase | 0.0923 | 1.1 | 0.085125 |
| YGL139W | FLC3 | unknown | 0.0923 | 1.1 | 0.581172 |
| YMR239C | RNT1 | ribonuclease III | 0.0927 | 1.1 | 0.762863 |
| YKR008W | RSC4 | unknown | 0.0927 | 1.1 | 0.637762 |
| YML113W | DAT1 | AT DNA binding | 0.0927 | 1.1 | 0.317989 |
| YLR201C | COQ9 | unknown | 0.0930 | 1.1 | 0.238061 |
| YJR011C | | unknown | 0.0935 | 1.1 | 0.454217 |
| YPL220W | RPL1A | structural constituent of ribosome | 0.0937 | 1.1 | 0.307226 |
| YIL050W | PCL7 | cyclin-dependent protein kinase | 0.0938 | 1.1 | 0.566823 |
| YOL072W | THP1 | protein binding | 0.0938 | 1.1 | 0.370865 |
| YGL066W | SGF73 | structural molecule * | 0.0940 | 1.1 | 0.036297 |
| YPL152W | RRD2 | protein phosphatase type 2A regulator | 0.0940 | 1.1 | 0.120268 |
| YLR068W | FYV7 | unknown | 0.0942 | 1.1 | 0.704986 |
| YDL236W | PHO13 | alkaline phosphatase * | 0.0942 | 1.1 | 0.170029 |
| YOL012C | HTZ1 | chromatin binding | 0.0948 | 1.1 | 0.283158 |
| YDR356W | SPC110 | structural constituent of cytoskeleton | 0.0950 | 1.1 | 0.580334 |
| YDR427W | RPN9 | structural molecule | 0.0950 | 1.1 | 0.381451 |
| YER013W | PRP22 | RNA splicing factor | 0.0952 | 1.1 | 0.522285 |
| YBR278W | DPB3 | epsilon DNA polymerase | 0.0952 | 1.1 | 0.66785 |
| YJL004C | SYS1 | unknown | 0.0955 | 1.1 | 0.759446 |
| YBR250W | | unknown | 0.0955 | 1.1 | 0.03594 |
| YMR133W | REC114 | unknown | 0.0960 | 1.1 | 0.544977 |
| Q0045 | COX1 | cytochrome-c oxidase | 0.0960 | 1.1 | 0.637676 |
| YCR091W | KIN82 | protein kinase | 0.0962 | 1.1 | 0.493797 |
| YKR029C | SET3 | NAD-dependent histone deacetylase * | 0.0963 | 1.1 | 0.339699 |
| YNR066C | | unknown | 0.0963 | 1.1 | 0.652778 |
| YLR310C | CDC25 | Ras guanyl-nucleotide exchange factor | 0.0963 | 1.1 | 0.0336884 |
| YPL133C | RDS2 | transcription factor | 0.0965 | 1.1 | 0.640981 |
| YML014W | TRM9 | tRNA (uridine) methyltransferase | 0.0967 | 1.1 | 0.161281 |
| YHR169W | DBP8 | ATP-dependent RNA helicase | 0.0970 | 1.1 | 0.115755 |
| YLR003C | | unknown | 0.0972 | 1.1 | 0.697762 |
| YMR282C | AEP2 | unknown | 0.0975 | 1.1 | 0.574056 |
| YPR048W | TAH18 | unknown | 0.0975 | 1.1 | 0.561827 |
| YGL189C | RPS26A | structural constituent of ribosome | 0.0977 | 1.1 | 0.49197 |
| YJR105W | ADO1 | adenosine kinase | 0.0977 | 1.1 | 0.468926 |
| YMR291W | | protein kinase | 0.0977 | 1.1 | 0.639081 |
| YDR074W | TPS2 | trehalose-phosphatase | 0.0980 | 1.1 | 0.126764 |
| YBR301W | DAN3 | unknown | 0.0980 | 1.1 | 0.650155 |
| YPL208W | RKM1 | protein-lysine N-methyltransferase | 0.0980 | 1.1 | 0.334358 |
| YKL121W | | unknown | 0.0982 | 1.1 | 0.46047 |
| YDL211C | | unknown | 0.0983 | 1.1 | 0.601651 |
| YDR531W | | pantothenate kinase | 0.0985 | 1.1 | 0.195369 |
| YGR294W | PAU12 | unknown | 0.0985 | 1.1 | 0.130403 |
| YFL027C | GYP8 | Rab GTPase activator | 0.0987 | 1.1 | 0.607376 |
| YHR034C | LYS1 | saccharopine dehydrogenase | 0.0988 | 1.1 | 0.58791 |
| YBR029C | CDS1 | phosphatidate cytidyltransferase | 0.0988 | 1.1 | 0.0414334 |
| YFL038C | YPT1 | GTPase | 0.0988 | 1.1 | 0.513632 |
| YPR099C | | | 0.0990 | 1.1 | 0.575795 |
| YMR186W | HSC82 | unfolded protein binding | 0.0992 | 1.1 | 0.274954 |
| YOR195W | SLK19 | unknown | 0.0992 | 1.1 | 0.833194 |
| YBL034C | STU1 | structural constituent of cytoskeleton | 0.0992 | 1.1 | 0.844424 |
| YDL104C | QRI7 | metalloendopeptidase | 0.0995 | 1.1 | 0.204384 |
| YMR134W | | unknown | 0.0997 | 1.1 | 0.367815 |
| YPL178W | CBC2 | RNA cap binding | 0.1003 | 1.1 | 0.0571149 |
| YFL028C | CAF16 | ATPase | 0.1005 | 1.1 | 0.736906 |
| YDR164C | SEC1 | SNARE binding | 0.1005 | 1.1 | 0.557822 |
| YGR216C | GPI1 | UDP-glycosyltransferase | 0.1007 | 1.1 | 0.50966 |
| YOL006C | TOP1 | DNA topoisomerase type I | 0.1007 | 1.1 | 0.405039 |
| YPL143W | RPL33A | structural constituent of ribosome | 0.1012 | 1.1 | 0.435448 |
| YLR273C | PIG1 | protein phosphatase type 1 regulator | 0.1015 | 1.1 | 0.310524 |
| YBL007C | SLA1 | protein binding | 0.1017 | 1.1 | 0.563892 |
| YBR089W | | | 0.1020 | 1.1 | 0.0155459 |
| YMR223W | UBP8 | ubiquitin-specific protease | 0.1025 | 1.1 | 0.673637 |
| YLR279W | | | 0.1030 | 1.1 | 0.574652 |
| YAR027W | UIP3 | unknown | 0.1032 | 1.1 | 0.131823 |
| YHR098C | SFB3 | unknown | 0.1032 | 1.1 | 0.193573 |
| YNL087W | TCB2 | unknown | 0.1033 | 1.1 | 0.630482 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YHR161C | YAP1801 | clathrin binding | 0.1037 | 1.1 | 0.290656 |
| YHL011C | PRS3 | ribose phosphate diphosphokinase | 0.1037 | 1.1 | 0.0660632 |
| YFR042W | | unknown | 0.1037 | 1.1 | 0.766893 |
| YJL093C | TOK1 | potassium channel | 0.1038 | 1.1 | 0.10765 |
| YBR113W | | | 0.1038 | 1.1 | 0.181183 |
| YPL153C | RAD53 | protein threonine/tyrosine kinase | 0.1040 | 1.1 | 0.725589 |
| YOR009W | TIR4 | unknown | 0.1042 | 1.1 | 0.648487 |
| YNL080C | | unknown | 0.1047 | 1.1 | 0.100427 |
| YBR195C | MSI1 | transcription regulator | 0.1047 | 1.1 | 0.485527 |
| YKL180W | RPL17A | structural constituent of ribosome | 0.1048 | 1.1 | 0.112591 |
| YGR150C | | unknown | 0.1050 | 1.1 | 0.562377 |
| YPR152C | URN1 | unknown | 0.1052 | 1.1 | 0.529878 |
| YMR091C | NPL6 | unknown | 0.1053 | 1.1 | 0.016653 |
| YOL088C | MPD2 | protein disulfide isomerase | 0.1055 | 1.1 | 0.275261 |
| YLR147C | SMD3 | RNA splicing factor | 0.1055 | 1.1 | 0.0530108 |
| YDR298C | ATP5 | structural molecule * | 0.1055 | 1.1 | 0.570317 |
| YER034W | | unknown | 0.1057 | 1.1 | 0.417125 |
| YFR028C | CDC14 | phosphoprotein phosphatase | 0.1060 | 1.1 | 0.823634 |
| YDR352W | | unknown | 0.1060 | 1.1 | 0.395957 |
| YBR152W | SPP381 | protein binding | 0.1060 | 1.1 | 0.656622 |
| YBR283C | SSH1 | protein transporter | 0.1060 | 1.1 | 0.669242 |
| YDL007W | RPT2 | ATPase * | 0.1062 | 1.1 | 0.0547079 |
| YKR091W | SRL3 | unknown | 0.1065 | 1.1 | 0.181936 |
| YNL096C | RPS7B | structural constituent of ribosome | 0.1072 | 1.1 | 0.0348401 |
| YLR398C | SKI2 | RNA helicase | 0.1077 | 1.1 | 0.672459 |
| YNR021W | | unknown | 0.1078 | 1.1 | 0.0966738 |
| YMR161W | HLJ1 | chaperone regulator | 0.1080 | 1.1 | 0.385789 |
| YPR107C | YTH1 | RNA binding | 0.1080 | 1.1 | 0.47814 |
| YGR041W | BUD9 | unknown | 0.1080 | 1.1 | 0.733979 |
| YMR212C | EFR3 | unknown | 0.1082 | 1.1 | 0.46249 |
| YDL081C | RPP1A | structural constituent of ribosome | 0.1083 | 1.1 | 0.139958 |
| YKL165C | MCD4 | unknown | 0.1083 | 1.1 | 0.744045 |
| YOR323C | PRO2 | glutamate-5-semialdehyde dehydrogenase | 0.1083 | 1.1 | 0.404042 |
| YOR275C | RIM20 | unknown | 0.1090 | 1.1 | 0.518349 |
| YIL006W | YIA6 | transporter | 0.1090 | 1.1 | 0.321675 |
| YLR272C | YCS4 | unknown | 0.1092 | 1.1 | 0.262957 |
| YCR059C | YIH1 | unknown | 0.1095 | 1.1 | 0.296258 |
| YNL133C | FYV6 | unknown | 0.1100 | 1.1 | 0.0698388 |
| YER041W | YEN1 | single-stranded DNA specific endodeoxyribonuclease | 0.1103 | 1.1 | 0.444506 |
| YGL100W | SEH1 | structural molecule | 0.1108 | 1.1 | 0.659628 |
| YGL044C | RNA15 | RNA binding | 0.1110 | 1.1 | 0.440088 |
| YGL131C | SNT2 | DNA binding | 0.1115 | 1.1 | 0.728825 |
| YDL051W | LHP1 | RNA binding | 0.1115 | 1.1 | 0.60111 |
| YKR075C | | unknown | 0.1115 | 1.1 | 0.268586 |
| YDL009C | | | 0.1117 | 1.1 | 0.656888 |
| YMR277W | FCP1 | phosphoprotein phosphatase * | 0.1117 | 1.1 | 0.338533 |
| YBR154C | RPB5 | DNA-directed RNA polymerase | 0.1118 | 1.1 | 0.501093 |
| YLR280C | | | 0.1120 | 1.1 | 0.202865 |
| YPL267W | ACM1 | unknown | 0.1122 | 1.1 | 0.661205 |
| YPL094C | SEC62 | protein binding | 0.1122 | 1.1 | 0.631389 |
| YOL013C | HRD1 | ubiquitin-protein ligase | 0.1128 | 1.1 | 0.258275 |
| YHR082C | KSP1 | protein serine/threonine kinase | 0.1128 | 1.1 | 0.783873 |
| YOL054W | PSH1 | unknown | 0.1128 | 1.1 | 0.215513 |
| YGR239C | PEX21 | protein binding | 0.1130 | 1.1 | 0.72442 |
| YGR280C | PXR1 | unknown | 0.1132 | 1.1 | 0.574029 |
| YNL121C | TOM70 | protein transporter | 0.1132 | 1.1 | 0.367403 |
| YJL126W | NIT2 | hydrolase , acting on carbon-nitrogen (but not peptide) bond | 0.1133 | 1.1 | 0.554981 |
| YDR262W | | unknown | 0.1138 | 1.1 | 0.198983 |
| YML104C | MDM1 | structural constituent of cytoskeleton | 0.1138 | 1.1 | 0.785894 |
| YDL153C | SAS10 | snoRNA binding | 0.1142 | 1.1 | 0.512859 |
| YPL063W | TIM50 | unknown | 0.1143 | 1.1 | 0.545651 |
| YHR108W | GGA2 | ubiquitin binding | 0.1145 | 1.1 | 0.572836 |
| YDR188W | CCT6 | unfolded protein binding | 0.1145 | 1.1 | 0.49423 |
| YKL098W | | unknown | 0.1145 | 1.1 | 0.402327 |
| YBL032W | HEK2 | mRNA binding | 0.1147 | 1.1 | 0.562439 |
| YER123W | YCK3 | casein kinase | 0.1148 | 1.1 | 0.146617 |
| YLR199C | | unknown | 0.1148 | 1.1 | 0.070085 |
| YJR128W | | | 0.1148 | 1.1 | 0.537542 |
| YAL035W | FUN12 | GTPase * | 0.1153 | 1.1 | 0.109083 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|--|----------------------------|-------------|------------|
| YCL054W | SPB1 | rRNA (uridine-2'-O-)-methyltransferase * | 0.1155 | 1.1 | 0.503113 |
| YKL004W | AUR1 | inositol phosphoceramide synthase | 0.1158 | 1.1 | 0.561091 |
| YBR258C | SHG1 | histone lysine N-methyltransferase (H3-K4 specific) | 0.1163 | 1.1 | 0.100662 |
| YMR208W | ERG12 | mevalonate kinase | 0.1165 | 1.1 | 0.216129 |
| YML095C | RAD10 | single-stranded DNA binding | 0.1167 | 1.1 | 0.101069 |
| YKR083C | DAD2 | structural constituent of cytoskeleton | 0.1167 | 1.1 | 0.0839237 |
| YBL101W-B | | | 0.1167 | 1.1 | 0.504111 |
| YNR003C | RPC34 | DNA-directed RNA polymerase | 0.1170 | 1.1 | 0.0551875 |
| YDL082W | RPL13A | structural constituent of ribosome | 0.1173 | 1.1 | 0.267745 |
| YLR154C | RNH203 | ribonuclease H | 0.1175 | 1.1 | 0.3495 |
| YPR086W | SUA7 | general RNA polymerase II transcription factor | 0.1180 | 1.1 | 0.521669 |
| YGL001C | ERG26 | C-3 sterol dehydrogenase | 0.1180 | 1.1 | 0.293019 |
| YOR280C | FSH3 | serine hydrolase | 0.1183 | 1.1 | 0.256269 |
| YDR340W | | | 0.1192 | 1.1 | 0.273447 |
| YDR136C | | | 0.1193 | 1.1 | 0.591126 |
| YLR333C | RPS25B | structural constituent of ribosome | 0.1193 | 1.1 | 0.284028 |
| YLR313C | SPH1 | cytoskeletal regulator | 0.1195 | 1.1 | 0.448867 |
| YDR449C | UTP6 | snoRNA binding | 0.1198 | 1.1 | 0.388719 |
| YOL051W | GAL11 | RNA polymerase II transcription mediator | 0.1200 | 1.1 | 0.606222 |
| YJL128C | PBS2 | MAP kinase | 0.1202 | 1.1 | 0.628837 |
| YOR157C | PUP1 | endopeptidase | 0.1205 | 1.1 | 0.11702 |
| YFL037W | TUB2 | structural constituent of cytoskeleton | 0.1208 | 1.1 | 0.345192 |
| YHR166C | CDC23 | protein binding | 0.1210 | 1.1 | 0.185775 |
| YOL084W | PHM7 | unknown | 0.1212 | 1.1 | 0.662521 |
| YPR145W | ASN1 | asparagine synthase (glutamine-hydrolyzing) | 0.1213 | 1.1 | 0.802176 |
| YDL129W | unknown | | 0.1218 | 1.1 | 0.60064 |
| YER128W | unknown | | 0.1223 | 1.1 | 0.00573564 |
| YAL032C | PRP45 | transcriptional activator | 0.1225 | 1.1 | 0.310836 |
| YLR191W | PEX13 | protein binding | 0.1235 | 1.1 | 0.485826 |
| YDL074C | BRE1 | ubiquitin-protein ligase | 0.1238 | 1.1 | 0.48786 |
| YPL140C | MKK2 | MAP kinase | 0.1240 | 1.1 | 0.688968 |
| YHR177W | unknown | | 0.1243 | 1.1 | 0.360452 |
| YLR210W | CLB4 | cyclin-dependent protein kinase | 0.1247 | 1.1 | 0.076248 |
| YEL053C | MAK10 | peptide alpha-N-acetyltransferase | 0.1250 | 1.1 | 0.185465 |
| YPL239W | YAR1 | unknown | 0.1250 | 1.1 | 0.162251 |
| YBL046W | PSY4 | unknown | 0.1252 | 1.1 | 0.425641 |
| YDL031W | DBP10 | ATP-dependent RNA helicase | 0.1258 | 1.1 | 0.173824 |
| YNL157W | IGO1 | unknown | 0.1258 | 1.1 | 0.152728 |
| YBR003W | COQ1 | trans-hexaprenyltranstransferase | 0.1260 | 1.1 | 0.506417 |
| YIL007C | NAS2 | unknown | 0.1265 | 1.1 | 0.315869 |
| YLR228C | ECM22 | RNA polymerase II transcription factor | 0.1267 | 1.1 | 0.24262 |
| YDL190C | UFD2 | ubiquitin conjugating enzyme | 0.1268 | 1.1 | 0.0742965 |
| YJL002C | OST1 | dolichyl-diphosphooligosaccharide-protein glycotransferase | 0.1277 | 1.1 | 0.517673 |
| YIL169C | unknown | | 0.1282 | 1.1 | 0.600069 |
| YBL033C | RIB1 | cyclohydrolase | 0.1283 | 1.1 | 0.0602846 |
| YBR111C | YSA1 | phosphoribosyl-ATP diphosphatase | 0.1285 | 1.1 | 0.123213 |
| YOL158C | ENB1 | ferric-enterobactin transporter | 0.1287 | 1.1 | 0.491319 |
| YDL137W | ARF2 | GTPase | 0.1288 | 1.1 | 0.0891944 |
| YGL122C | NAB2 | poly(A) binding | 0.1290 | 1.1 | 0.338914 |
| YLR317W | | | 0.1292 | 1.1 | 0.0329692 |
| YKR079C | TRZ1 | purine nucleotide binding | 0.1292 | 1.1 | 0.264391 |
| YMR304C-A | | | 0.1295 | 1.1 | 0.137957 |
| YBL081W | unknown | | 0.1297 | 1.1 | 0.194578 |
| YJR030C | unknown | | 0.1298 | 1.1 | 0.0414806 |
| YLL062C | MHT1 | homocysteine S-methyltransferase | 0.1303 | 1.1 | 0.644286 |
| YGL018C | JAC1 | chaperone binding | 0.1308 | 1.1 | 0.729041 |
| YJR031C | GEA1 | ARF guanyl-nucleotide exchange factor | 0.1308 | 1.1 | 0.569886 |
| YGL221C | NIF3 | unknown | 0.1308 | 1.1 | 0.147054 |
| YLR440C | SEC39 | unknown | 0.1310 | 1.1 | 0.0588094 |
| YEL018W | EAF5 | unknown | 0.1312 | 1.1 | 0.605228 |
| YDR269C | | | 0.1313 | 1.1 | 0.193176 |
| YFR018C | unknown | | 0.1313 | 1.1 | 0.110047 |
| YMR044W | IOC4 | protein binding | 0.1315 | 1.1 | 0.630978 |
| YLR360W | VPS38 | unknown | 0.1317 | 1.1 | 0.153893 |
| YFL042C(1) | | | 0.1317 | 1.1 | 0.154369 |
| YHR005C | GPA1 | GTPase | 0.1318 | 1.1 | 0.424583 |
| YDR100W | TVP15 | unknown | 0.1320 | 1.1 | 0.370396 |
| YBR164C | ARL1 | GTPase | 0.1327 | 1.1 | 0.0427965 |
| YLL013C | PUF3 | mRNA binding | 0.1327 | 1.1 | 0.208463 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|------------|
| YBR136W | MEC1 | protein kinase | 0.1327 | 1.1 | 0.478742 |
| YKL214C | YRA2 | RNA binding | 0.1328 | 1.1 | 0.546143 |
| YER189W | | unknown | 0.1330 | 1.1 | 0.559705 |
| YOR394W | PAU21 | unknown | 0.1330 | 1.1 | 0.364599 |
| YCL004W | PGS1 | CDP-diacylglycerol-glycerol-3-phosphate 3-phosphatidyltra | 0.1333 | 1.1 | 0.538722 |
| YDL030W | PRP9 | RNA binding | 0.1333 | 1.1 | 0.420907 |
| Q0065 | AI4 | endonuclease | 0.1335 | 1.1 | 0.550689 |
| YMR197C | VTI1 | v-SNARE | 0.1337 | 1.1 | 0.18774 |
| YHR135C | YCK1 | casein kinase | 0.1337 | 1.1 | 0.288841 |
| YDR443C | SSN2 | general RNA polymerase II transcription factor | 0.1338 | 1.1 | 0.0916875 |
| YEL037C | RAD23 | protein binding | 0.1340 | 1.1 | 0.358748 |
| YPR191W | QCR2 | ubiquinol-cytochrome-c reductase | 0.1340 | 1.1 | 0.118421 |
| YNL298W | CLA4 | protein serine/threonine kinase | 0.1345 | 1.1 | 0.595546 |
| YBL098W | BNA4 | kynurenine 3-monooxygenase | 0.1347 | 1.1 | 0.153824 |
| YOR059C | | unknown | 0.1350 | 1.1 | 0.0949166 |
| YAR007C | RFA1 | single-stranded DNA binding | 0.1352 | 1.1 | 0.20586 |
| YDL061C | RPS29B | structural constituent of ribosome | 0.1355 | 1.1 | 0.0872979 |
| YDR203W | | | 0.1360 | 1.1 | 0.182824 |
| YMR293C | | amidase | 0.1365 | 1.1 | 0.140582 |
| YAL068C | PAU8 | unknown | 0.1368 | 1.1 | 0.405266 |
| YCR020C | PET18 | unknown | 0.1368 | 1.1 | 0.224054 |
| YLL036C | PRP19 | RNA splicing factor | 0.1372 | 1.1 | 0.340738 |
| YLR323C | CWC24 | unknown | 0.1372 | 1.1 | 0.475044 |
| YER161C | SPT2 | DNA binding | 0.1375 | 1.1 | 0.139997 |
| YER075C | PTP3 | protein tyrosine phosphatase | 0.1377 | 1.1 | 0.0634348 |
| YML102C-A | | | 0.1378 | 1.1 | 0.528052 |
| YBR242W | | unknown | 0.1383 | 1.1 | 0.358679 |
| YHR067W | HTD2 | 3-hydroxyacyl-[acyl-carrier protein] dehydratase | 0.1385 | 1.1 | 0.711139 |
| YDR496C | PUF6 | specific transcriptional repressor | 0.1387 | 1.1 | 0.0849547 |
| YCR066W | RAD18 | ubiquitin conjugating enzyme * | 0.1392 | 1.1 | 0.336428 |
| YKR002W | PAP1 | polynucleotide adenylyltransferase | 0.1393 | 1.1 | 0.12771 |
| YJL162C | JJJ2 | unknown | 0.1398 | 1.1 | 0.256489 |
| YFR030W | MET10 | sulfite reductase (NADPH) | 0.1400 | 1.1 | 0.608248 |
| YDL183C | | unknown | 0.1400 | 1.1 | 0.338239 |
| YBR207W | FTH1 | iron ion transporter | 0.1402 | 1.1 | 0.272778 |
| YDR535C | | | 0.1407 | 1.1 | 0.451928 |
| YFL020C | PAU5 | unknown | 0.1407 | 1.1 | 0.512559 |
| YMR272C | SCS7 | oxidoreductase | 0.1410 | 1.1 | 0.541139 |
| YER061C | CEM1 | fatty-acid synthase | 0.1412 | 1.1 | 0.281065 |
| YLR359W | ADE13 | adenylosuccinate lyase | 0.1412 | 1.1 | 0.541508 |
| YOR106W | VAM3 | t-SNARE | 0.1413 | 1.1 | 0.508981 |
| YBL105C | PKC1 | protein kinase | 0.1413 | 1.1 | 0.376838 |
| YIL172C | | glucosidase | 0.1413 | 1.1 | 0.0936326 |
| YCR082W | AHC2 | unknown | 0.1420 | 1.1 | 0.232784 |
| YCR090C | | unknown | 0.1420 | 1.1 | 0.243995 |
| YDR088C | SLU7 | RNA splicing factor | 0.1423 | 1.1 | 0.399517 |
| YLR395C | COX8 | cytochrome-c oxidase | 0.1423 | 1.1 | 0.272628 |
| YMR175W | SIP18 | phospholipid binding | 0.1428 | 1.1 | 0.425012 |
| YBR233W | PBP2 | unknown | 0.1432 | 1.1 | 0.408402 |
| YGR034W | RPL26B | structural constituent of ribosome* | 0.1437 | 1.1 | 0.175154 |
| YML024W | RPS17A | structural constituent of ribosome | 0.1440 | 1.1 | 0.400271 |
| YDR077W | SED1 | structural constituent of cell wall | 0.1442 | 1.1 | 0.0491482 |
| YGR003W | CUL3 | protein binding | 0.1442 | 1.1 | 0.00275001 |
| YCR095C | OCA4 | unknown | 0.1442 | 1.1 | 0.151949 |
| YMR237W | BCH1 | unknown | 0.1445 | 1.1 | 0.290032 |
| YNL084C | END3 | protein binding | 0.1447 | 1.1 | 0.15514 |
| YDR306C | | protein binding | 0.1450 | 1.1 | 0.0659009 |
| YEL059W | | | 0.1450 | 1.1 | 0.650518 |
| YGR212W | SLI1 | N-acetyltransferase | 0.1452 | 1.1 | 0.704562 |
| YDL159W | STE7 | MAP kinase | 0.1453 | 1.1 | 0.485021 |
| YNL069C | RPL16B | structural constituent of ribosome* | 0.1453 | 1.1 | 0.176655 |
| YKR053C | YSR3 | sphingosine-1-phosphate phosphatase | 0.1455 | 1.1 | 0.445431 |
| YCR044C | PER1 | unknown | 0.1457 | 1.1 | 0.032751 |
| YDR090C | | unknown | 0.1457 | 1.1 | 0.228858 |
| YDR453C | TSA2 | thioredoxin peroxidase | 0.1463 | 1.1 | 0.216189 |
| YDR287W | | inositol-1(or 4)-monophosphatase | 0.1463 | 1.1 | 0.456887 |
| YBR035C | PDX3 | pyridoxamine-phosphate oxidase | 0.1465 | 1.1 | 0.205745 |
| YDL001W | RMD1 | unknown | 0.1467 | 1.1 | 0.491882 |
| YNL094W | APP1 | unknown | 0.1470 | 1.1 | 0.481346 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|--|----------------------------|-------------|------------|
| YOR126C | IAH1 | hydrolase , acting on ester bonds | 0.1470 | 1.1 | 0.180527 |
| YER019W | ISC1 | phospholipase C | 0.1470 | 1.1 | 0.360717 |
| YCL014W(1) | | | 0.1475 | 1.1 | 0.599441 |
| YOR091W | TMA46 | unknown | 0.1478 | 1.1 | 0.016444 |
| YDR465C | RMT2 | protein-arginine N-methyltransferase | 0.1480 | 1.1 | 0.512471 |
| YCR014C | POL4 | beta DNA polymerase | 0.1482 | 1.1 | 0.178999 |
| YGR042W | | unknown | 0.1487 | 1.1 | 0.40785 |
| YMR014W | BUD22 | unknown | 0.1488 | 1.1 | 0.594856 |
| YLR448W | RPL6B | structural constituent of ribosome* | 0.1492 | 1.1 | 0.285884 |
| YLR443W | ECM7 | unknown | 0.1493 | 1.1 | 0.744632 |
| YKL054C | DEF1 | unknown | 0.1493 | 1.1 | 0.304704 |
| YGL236C | MT01 | unknown | 0.1493 | 1.1 | 0.44651 |
| YNR052C | POP2 | 3'-5'-exoribonuclease | 0.1495 | 1.1 | 0.0901193 |
| YBL052C | SAS3 | histone acetyltransferase * | 0.1497 | 1.1 | 0.62486 |
| YLR238W | FAR10 | unknown | 0.1498 | 1.1 | 0.275035 |
| YEL004W | YEA4 | UDP-N-acetylglucosamine transporter | 0.1500 | 1.1 | 0.375419 |
| YKR022C | NTR2 | unknown | 0.1503 | 1.1 | 0.514403 |
| YNL149C | PGA2 | unknown | 0.1507 | 1.1 | 0.00900468 |
| YDR456W | NHX1 | monovalent inorganic cation transporter | 0.1508 | 1.1 | 0.381317 |
| YHL038C | CBP2 | RNA splicing factor | 0.1508 | 1.1 | 0.484607 |
| YLR240W | VPS34 | protein kinase | 0.1515 | 1.1 | 0.151736 |
| YIL104C | SHQ1 | unknown | 0.1517 | 1.1 | 0.536381 |
| YJR068W | RFC2 | DNA clamp loader * | 0.1518 | 1.1 | 0.285315 |
| YPR124W | CTR1 | copper uptake transporter | 0.1520 | 1.1 | 0.643434 |
| YNL152W | | phospholipid binding | 0.1523 | 1.1 | 0.584724 |
| YKR092C | SRP40 | unfolded protein binding | 0.1525 | 1.1 | 0.373049 |
| YPR037C | ERV2 | thiol oxidase | 0.1527 | 1.1 | 0.379644 |
| YDL072C | YET3 | unknown | 0.1528 | 1.1 | 0.423548 |
| YKL007W | CAP1 | actin filament binding | 0.1528 | 1.1 | 0.345026 |
| YTR018W | YAP5 | RNA polymerase II transcription factor | 0.1528 | 1.1 | 0.289688 |
| YDR243C | PRP28 | RNA splicing factor | 0.1528 | 1.1 | 0.0768341 |
| YML119W | | unknown | 0.1530 | 1.1 | 0.284889 |
| YPR061C | JID1 | unknown | 0.1530 | 1.1 | 0.0236308 |
| YBL036C | | alanine racemase * | 0.1532 | 1.1 | 0.357397 |
| YER021W | RPN3 | unknown | 0.1532 | 1.1 | 0.1174 |
| YDR361C | BCP1 | unknown | 0.1537 | 1.1 | 0.0211505 |
| YBR002C | RER2 | prenyltransferase * | 0.1538 | 1.1 | 0.725365 |
| YDR310C | SUM1 | transcriptional repressor | 0.1548 | 1.1 | 0.37048 |
| YOR239W | ABP140 | S-adenosylmethionine-dependent methyltransferase * | 0.1548 | 1.1 | 0.312529 |
| YOR025W | HST3 | DNA binding | 0.1548 | 1.1 | 0.301654 |
| YKL019W | RAM2 | protein farnesyltransferase * | 0.1550 | 1.1 | 0.297034 |
| YCL048W | SPS22 | unknown | 0.1553 | 1.1 | 0.503054 |
| YER001W | MNN1 | alpha-1,3-mannosyltransferase | 0.1553 | 1.1 | 0.103183 |
| YEL076C-A | | unknown | 0.1557 | 1.1 | 0.221497 |
| YNL206C | RTT106 | unknown | 0.1558 | 1.1 | 0.169254 |
| YGR040W | KSS1 | MAP kinase | 0.1560 | 1.1 | 0.216823 |
| YGR240C | PFK1 | 6-phosphofructokinase | 0.1562 | 1.1 | 0.11625 |
| YDL012C | | unknown | 0.1567 | 1.1 | 0.081968 |
| YLR065C | | unknown | 0.1568 | 1.1 | 0.257181 |
| YMR075W | RCO1 | unknown | 0.1572 | 1.1 | 0.160801 |
| YBR128C | ATG14 | unknown | 0.1575 | 1.1 | 0.326089 |
| YOR288C | MPD1 | protein disulfide isomerase | 0.1575 | 1.1 | 0.589195 |
| YLR435W | TSR2 | unknown | 0.1578 | 1.1 | 0.0166818 |
| YER107C | GLE2 | structural molecule | 0.1582 | 1.1 | 0.414265 |
| YDL067C | COX9 | cytochrome-c oxidase | 0.1590 | 1.1 | 0.225618 |
| YKL216W | URA1 | dihydroorotate dehydrogenase | 0.1590 | 1.1 | 0.359024 |
| YLR299W | ECM38 | protein-glutamine gamma-glutamyltransferase | 0.1592 | 1.1 | 0.0204593 |
| YMR144W | | unknown | 0.1598 | 1.1 | 0.222308 |
| YIL175W | | unknown | 0.1598 | 1.1 | 0.57123 |
| YER043C | SAH1 | adenosylhomocysteinase | 0.1602 | 1.1 | 0.228393 |
| YHR195W | NVJ1 | protein binding | 0.1602 | 1.1 | 0.166609 |
| YER014W | HEM14 | protoporphyrinogen oxidase | 0.1603 | 1.1 | 0.116793 |
| YDR311W | TFB1 | general RNA polymerase II transcription factor | 0.1603 | 1.1 | 0.134742 |
| YKL013C | ARC19 | structural molecule | 0.1607 | 1.1 | 0.191698 |
| YBR274W | CHK1 | protein kinase | 0.1608 | 1.1 | 0.11033 |
| YLR145W | RMP1 | unknown | 0.1610 | 1.1 | 0.36634 |
| YGR203W | | protein tyrosine phosphatase * | 0.1613 | 1.1 | 0.203448 |
| YHR076W | PTC7 | protein phosphatase type 2C | 0.1617 | 1.1 | 0.239947 |
| YDL233W | | unknown | 0.1617 | 1.1 | 0.0213597 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YER066W | | unknown | 0.1618 | 1.1 | 0.416982 |
| YLR196W | PWP1 | unknown | 0.1620 | 1.1 | 0.397123 |
| YOL052C | SPE2 | adenosylmethionine decarboxylase | 0.1622 | 1.1 | 0.11487 |
| YPL210C | SRP72 | signal sequence binding | 0.1623 | 1.1 | 0.634838 |
| YIR023W | DAL81 | specific RNA polymerase II transcription factor | 0.1625 | 1.1 | 0.37432 |
| YIL077C | | unknown | 0.1627 | 1.1 | 0.22355 |
| YBR249C | ARO4 | 3-deoxy-7-phosphoheptulonate synthase | 0.1633 | 1.1 | 0.301179 |
| YLR405W | DUS4 | tRNA dihydrouridine synthase | 0.1633 | 1.1 | 0.405922 |
| YBR191W | RPL21A | structural constituent of ribosome | 0.1633 | 1.1 | 0.320859 |
| YOR308C | SNU66 | RNA splicing factor | 0.1635 | 1.1 | 0.403565 |
| YGR201C | | unknown | 0.1647 | 1.1 | 0.501478 |
| YDR444W | | unknown | 0.1648 | 1.1 | 0.397247 |
| YMR097C | MTG1 | GTPase | 0.1653 | 1.1 | 0.0950438 |
| YNR018W | | unknown | 0.1653 | 1.1 | 0.0247463 |
| YOL004W | SIN3 | histone deacetylase | 0.1660 | 1.1 | 0.409466 |
| YDL056W | MBP1 | DNA binding | 0.1662 | 1.1 | 0.412157 |
| YER057C | HMF1 | unknown | 0.1662 | 1.1 | 0.29538 |
| YIL001W | | unknown | 0.1665 | 1.1 | 0.612778 |
| YJL223C | PAU1 | unknown | 0.1667 | 1.1 | 0.169025 |
| YER122C | GLO3 | ARF GTPase activator | 0.1670 | 1.1 | 0.458502 |
| YNL264C | PDR17 | phosphatidylinositol transporter | 0.1670 | 1.1 | 0.0993254 |
| YJL136C | RPS21B | structural constituent of ribosome | 0.1672 | 1.1 | 0.162568 |
| YPL099C | FMP14 | unknown | 0.1675 | 1.1 | 0.491244 |
| YOR113W | AZF1 | DNA binding | 0.1675 | 1.1 | 0.053153 |
| YPL088W | | aryl-alcohol dehydrogenase | 0.1675 | 1.1 | 0.130449 |
| YGL071W | RCS1 | transcription factor | 0.1677 | 1.1 | 0.477391 |
| YIL065C | FIS1 | unknown | 0.1678 | 1.1 | 0.176653 |
| YHR199C | FMP34 | unknown | 0.1680 | 1.1 | 0.117084 |
| YPR187W | RPO26 | DNA-directed RNA polymerase | 0.1685 | 1.1 | 0.204089 |
| YCR105W | ADH7 | alcohol dehydrogenase | 0.1687 | 1.1 | 0.0830892 |
| YIL043C | CBR1 | cytochrome-b5 reductase | 0.1688 | 1.1 | 0.0367232 |
| YMR192W | GYL1 | protein binding | 0.1690 | 1.1 | 0.40146 |
| YGR101W | PCP1 | peptidase | 0.1692 | 1.1 | 0.50063 |
| YPR072W | NOT5 | 3'-5'-exoribonuclease | 0.1692 | 1.1 | 0.58087 |
| YLR261C | | | 0.1697 | 1.1 | 0.636401 |
| YIL039W | | unknown | 0.1700 | 1.1 | 0.0329626 |
| YHR017W | YSC83 | unknown | 0.1702 | 1.1 | 0.0372833 |
| YMR052W | FAR3 | unknown | 0.1703 | 1.1 | 0.542139 |
| YPL059W | GRX5 | thiol-disulfide exchange intermediate | 0.1703 | 1.1 | 0.320115 |
| YDR279W | RNH202 | ribonuclease H | 0.1705 | 1.1 | 0.447241 |
| YBR107C | IML3 | unknown | 0.1705 | 1.1 | 0.602537 |
| YOR003W | YSP3 | peptidase | 0.1708 | 1.1 | 0.298875 |
| YIL019W | FAF1 | unknown | 0.1708 | 1.1 | 0.28448 |
| YGR128C | UTP8 | snoRNA binding | 0.1710 | 1.1 | 0.0728068 |
| YDR327W | | | 0.1710 | 1.1 | 0.147889 |
| YDL243C | AAD4 | aryl-alcohol dehydrogenase | 0.1712 | 1.1 | 0.225989 |
| YNL003C | PET8 | S-adenosylmethionine transporter | 0.1712 | 1.1 | 0.337357 |
| YPR106W | ISR1 | protein kinase | 0.1715 | 1.1 | 0.252978 |
| YDR152W | GIR2 | unknown | 0.1715 | 1.1 | 0.380794 |
| YNL011C | | unknown | 0.1715 | 1.1 | 0.231544 |
| YEL064C | AVT2 | transporter | 0.1717 | 1.1 | 0.446737 |
| YOL030W | GAS5 | 1,3-beta-glucanosyltransferase | 0.1717 | 1.1 | 0.177334 |
| YLR382C | NAM2 | mRNA binding | 0.1722 | 1.1 | 0.149685 |
| YLL001W | DNM1 | GTPase | 0.1728 | 1.1 | 0.109521 |
| YDL099W | BUG1 | unknown | 0.1730 | 1.1 | 0.294758 |
| YML050W | | unknown | 0.1732 | 1.1 | 0.356024 |
| YOR391C | HSP33 | unfolded protein binding | 0.1735 | 1.1 | 0.156141 |
| YBR166C | TYR1 | prephenate dehydrogenase | 0.1737 | 1.1 | 0.0193129 |
| YBR259W | | unknown | 0.1747 | 1.1 | 0.23908 |
| YDR454C | GUK1 | guanylate kinase | 0.1747 | 1.1 | 0.209716 |
| YGL097W | SRM1 | signal transducer | 0.1747 | 1.1 | 0.368711 |
| YAR003W | SWD1 | histone lysine N-methyltransferase (H3-K4 specific) | 0.1752 | 1.1 | 0.23003 |
| YPL101W | ELP4 | RNA polymerase II transcription elongation factor | 0.1753 | 1.1 | 0.344681 |
| YDL143W | CCT4 | unfolded protein binding | 0.1755 | 1.1 | 0.544732 |
| YKL175W | ZRT3 | zinc ion transporter | 0.1755 | 1.1 | 0.423098 |
| YDL181W | INH1 | enzyme inhibitor | 0.1758 | 1.1 | 0.0270924 |
| YDR495C | VPS3 | unknown | 0.1762 | 1.1 | 0.336126 |
| YKL163W | PIR3 | structural constituent of cell wall | 0.1762 | 1.1 | 0.0809432 |
| YJL073W | JEM1 | unfolded protein binding | 0.1762 | 1.1 | 0.109618 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YMR166C | | transporter | 0.1762 | 1.1 | 0.465635 |
| YML038C | YMD8 | nucleotide-sugar transporter | 0.1765 | 1.1 | 0.405977 |
| YMR020W | FMS1 | amine oxidase | 0.1767 | 1.1 | 0.444654 |
| YDL206W | | unknown | 0.1767 | 1.1 | 0.0698194 |
| YDL146W | LDB17 | unknown | 0.1777 | 1.1 | 0.402632 |
| YML055W | SPC2 | protein binding | 0.1777 | 1.1 | 0.147956 |
| YML018C | | unknown | 0.1777 | 1.1 | 0.710491 |
| YGR014W | MSB2 | osmosensor | 0.1783 | 1.1 | 0.535058 |
| YPR062W | FCY1 | cytosine deaminase | 0.1785 | 1.1 | 0.135592 |
| YKL025C | PAN3 | poly(A)-specific ribonuclease | 0.1787 | 1.1 | 0.52356 |
| YJL165C | HAL5 | protein kinase | 0.1787 | 1.1 | 0.0704927 |
| YGR083C | GCD2 | translation initiation factor | 0.1790 | 1.1 | 0.632362 |
| YGR185C | TYS1 | tyrosine-tRNA ligase | 0.1790 | 1.1 | 0.101729 |
| YBR073W | RDH54 | DNA-dependent ATPase * | 0.1795 | 1.1 | 0.319776 |
| YNR032W | PPG1 | protein phosphatase type 2A | 0.1802 | 1.1 | 0.289962 |
| YOR261C | RPN8 | unknown | 0.1807 | 1.1 | 0.0713078 |
| YLR256W | HAP1 | specific RNA polymerase II transcription factor | 0.1808 | 1.1 | 0.103615 |
| YMR265C | | unknown | 0.1808 | 1.1 | 0.256657 |
| YLR092W | SUL2 | sulfate transporter | 0.1810 | 1.1 | 0.393825 |
| YBR048W | RPS11B | structural constituent of ribosome | 0.1813 | 1.1 | 0.168943 |
| YHR153C | SPO16 | unknown | 0.1813 | 1.1 | 0.432357 |
| YNL076W | MKS1 | transcriptional repressor | 0.1815 | 1.1 | 0.481046 |
| YHR031C | RRM3 | DNA helicase | 0.1818 | 1.1 | 0.335263 |
| YOR070C | GYP1 | Rab GTPase activator | 0.1823 | 1.1 | 0.0348458 |
| YNL027W | CRZ1 | transcription factor | 0.1823 | 1.1 | 0.0808546 |
| YKL018W | SWD2 | histone lysine N-methyltransferase (H3-K4 specific) | 0.1825 | 1.1 | 0.406908 |
| YIL064W | | S-adenosylmethionine-dependent methyltransferase | 0.1835 | 1.1 | 0.170483 |
| YHR133C | NSG1 | protein binding | 0.1837 | 1.1 | 0.105198 |
| YLR287C | | unknown | 0.1838 | 1.1 | 0.435417 |
| YDR376W | ARH1 | NADPH-adrenodoxin reductase | 0.1840 | 1.1 | 0.26118 |
| YPL183C | | unknown | 0.1843 | 1.1 | 0.572302 |
| YJL203W | PRP21 | RNA binding | 0.1845 | 1.1 | 0.416913 |
| YIL127C | | unknown | 0.1847 | 1.1 | 0.177489 |
| YML066C | SMA2 | unknown | 0.1848 | 1.1 | 0.681611 |
| YGR169C | PUS6 | pseudouridylyl synthase | 0.1848 | 1.1 | 0.558649 |
| YDL154W | MSH5 | unknown | 0.1850 | 1.1 | 0.0226532 |
| YDR215C | | | 0.1855 | 1.1 | 0.548794 |
| YCR058C | | | 0.1857 | 1.1 | 0.0256472 |
| YBR215W | HPC2 | transcription regulator | 0.1857 | 1.1 | 0.120199 |
| YPL034W | | unknown | 0.1860 | 1.1 | 0.772794 |
| YMR311C | GLC8 | enzyme activator | 0.1862 | 1.1 | 0.295748 |
| YMR322C | SNO4 | unfolded protein binding | 0.1862 | 1.1 | 0.432987 |
| YPL213W | LEA1 | RNA splicing factor | 0.1863 | 1.1 | 0.100498 |
| YMR098C | | unknown | 0.1865 | 1.1 | 0.265559 |
| YDR542W | PAU10 | unknown | 0.1867 | 1.1 | 0.262998 |
| YBR063C | | unknown | 0.1868 | 1.1 | 0.143337 |
| YBR285W | | unknown | 0.1872 | 1.1 | 0.0711402 |
| YOR171C | LCB4 | D-erythro-sphingosine kinase | 0.1872 | 1.1 | 0.180416 |
| YKR059W | TIF1 | translation initiation factor | 0.1875 | 1.1 | 0.166134 |
| YBR253W | SRB6 | RNA polymerase II transcription mediator | 0.1885 | 1.1 | 0.440593 |
| YKL189W | HYM1 | unknown | 0.1887 | 1.1 | 0.240389 |
| YDR212W | TCP1 | unfolded protein binding | 0.1887 | 1.1 | 0.553483 |
| YHR216W | IMD2 | IMP dehydrogenase | 0.1897 | 1.1 | 0.356608 |
| YDL207W | GLE1 | unknown | 0.1897 | 1.1 | 0.104677 |
| YNL160W | YGP1 | unknown | 0.1902 | 1.1 | 0.0411112 |
| YOR205C | FMP38 | unknown | 0.1905 | 1.1 | 0.0141179 |
| YDR423C | CAD1 | RNA polymerase II transcription factor | 0.1910 | 1.1 | 0.664048 |
| YNL287W | SEC21 | unknown | 0.1912 | 1.1 | 0.252102 |
| YER009W | NTF2 | Ran GTPase binding | 0.1912 | 1.1 | 0.314015 |
| YGR255C | COQ6 | ubiquinone biosynthesis monooxygenase | 0.1912 | 1.1 | 0.304234 |
| YGR286C | BIO2 | biotin synthase | 0.1915 | 1.1 | 0.155083 |
| YJR119C | | unknown | 0.1917 | 1.1 | 0.639129 |
| YNL090W | RHO2 | GTPase * | 0.1917 | 1.1 | 0.233709 |
| YGR198W | | transferase | 0.1918 | 1.1 | 0.197213 |
| YKL201C | MNN4 | unknown | 0.1923 | 1.1 | 0.626439 |
| YDR288W | NSE3 | DNA binding | 0.1935 | 1.1 | 0.177183 |
| YNL062C | GCD10 | tRNA binding | 0.1935 | 1.1 | 0.0723126 |
| YKR068C | BET3 | unknown | 0.1937 | 1.1 | 0.222498 |
| YML131W | | unknown | 0.1938 | 1.1 | 0.0895904 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YDR338C | | unknown | 0.1940 | 1.1 | 0.290121 |
| YBL099W | ATP1 | hydrogen-transporting ATP synthase , rotational mechanism | 0.1940 | 1.1 | 0.0538191 |
| YJL010C | NOP9 | RNA binding | 0.1943 | 1.1 | 0.163332 |
| YLR020C | YEH2 | sterol esterase | 0.1952 | 1.1 | 0.0119782 |
| YIL062C | ARC15 | structural molecule * | 0.1952 | 1.1 | 0.347166 |
| YNR049C | MSO1 | unknown | 0.1955 | 1.1 | 0.206031 |
| YBR077C | SLM4 | phosphatidylinositol-3,4-bisphosphate binding | 0.1957 | 1.1 | 0.142525 |
| YDR383C | NKP1 | unknown | 0.1958 | 1.1 | 0.262588 |
| YDR500C | RPL37B | structural constituent of ribosome | 0.1972 | 1.1 | 0.408698 |
| YDR365C | ESF1 | RNA binding | 0.1973 | 1.1 | 0.440112 |
| YAR002W | NUP60 | structural constituent of nuclear pore | 0.1975 | 1.1 | 0.00669003 |
| YPR199C | ARR1 | RNA polymerase II transcription factor | 0.1975 | 1.1 | 0.0310066 |
| YDR143C | SAN1 | ubiquitin-protein ligase | 0.1982 | 1.1 | 0.0396306 |
| YBR261C | | S-adenosylmethionine-dependent methyltransferase | 0.1983 | 1.1 | 0.0473883 |
| YCL036W | GFD2 | unknown | 0.1983 | 1.1 | 0.251213 |
| YPL206C | | glycerophosphodiester phosphodiesterase | 0.1985 | 1.1 | 0.0894852 |
| YOR120W | GCY1 | aldo-keto reductase | 0.1987 | 1.1 | 0.305555 |
| YDL102W | CDC2 | delta DNA polymerase * | 0.1988 | 1.1 | 0.355927 |
| YKR013W | PRY2 | unknown | 0.1990 | 1.1 | 0.501651 |
| YPR110C | RPC40 | DNA-directed RNA polymerase | 0.1992 | 1.1 | 0.157473 |
| YHR020W | | proline-tRNA ligase | 0.1992 | 1.1 | 0.312284 |
| YOR122C | PFY1 | actin monomer binding | 0.1995 | 1.1 | 0.0886395 |
| YBL080C | PET112 | unknown | 0.1998 | 1.1 | 0.278379 |
| YGL094C | PAN2 | poly(A)-specific ribonuclease | 0.2000 | 1.1 | 0.345 |
| YER171W | RAD3 | general RNA polymerase II transcription factor | 0.2003 | 1.1 | 0.482323 |
| YJR133W | XPT1 | xanthine phosphoribosyltransferase | 0.2005 | 1.1 | 0.0262605 |
| YMR070W | MOT3 | DNA binding | 0.2005 | 1.1 | 0.00594312 |
| YKL172W | EBP2 | unknown | 0.2007 | 1.1 | 0.187262 |
| YPL068C | | unknown | 0.2013 | 1.1 | 0.343503 |
| YIL105C | SLM1 | phosphoinositide binding | 0.2017 | 1.2 | 0.0844178 |
| YPL211W | NIP7 | unknown | 0.2017 | 1.2 | 0.13289 |
| YOR349W | CIN1 | beta-tubulin binding | 0.2018 | 1.2 | 0.611145 |
| YHL039W | | unknown | 0.2023 | 1.2 | 0.0691562 |
| YCR060W | TAH1 | chaperone binding | 0.2023 | 1.2 | 0.0452439 |
| YDR382W | RPP2B | structural constituent of ribosome | 0.2023 | 1.2 | 0.104016 |
| YOR166C | | unknown | 0.2025 | 1.2 | 0.228483 |
| YLL023C | | unknown | 0.2027 | 1.2 | 0.148691 |
| YCR094W | CDC50 | phospholipid-translocating ATPase | 0.2027 | 1.2 | 0.275728 |
| YIL157C | FMP35 | unknown | 0.2028 | 1.2 | 0.00442258 |
| YMR145C | NDE1 | NADH dehydrogenase | 0.2032 | 1.2 | 0.181511 |
| YIR030C | DCG1 | unknown | 0.2033 | 1.2 | 0.451877 |
| YGL191W | COX13 | enzyme regulator | 0.2033 | 1.2 | 0.148849 |
| YPR180W | AOS1 | SUMO activating enzyme | 0.2038 | 1.2 | 0.241394 |
| YCL027W | FUS1 | unknown | 0.2040 | 1.2 | 0.283427 |
| YAR073W | | | 0.2042 | 1.2 | 0.395782 |
| YNR045W | PET494 | translation regulator | 0.2042 | 1.2 | 0.211323 |
| YBR108W | | unknown | 0.2043 | 1.2 | 0.000884059 |
| YKL041W | VPS24 | protein binding | 0.2047 | 1.2 | 0.459616 |
| YLR110C | CCW12 | unknown | 0.2047 | 1.2 | 0.173838 |
| YCLX05C | | | 0.2052 | 1.2 | 0.386081 |
| YBL020W | RFT1 | oligosaccharide transporter | 0.2052 | 1.2 | 0.150556 |
| YMR128W | ECM16 | RNA helicase | 0.2057 | 1.2 | 0.0371025 |
| YMR244C-A | | unknown | 0.2057 | 1.2 | 0.0980733 |
| YDR480W | DIG2 | transcription factor | 0.2063 | 1.2 | 0.553812 |
| YMR081C | ISF1 | unknown | 0.2063 | 1.2 | 0.049114 |
| YLR185W | RPL37A | structural constituent of ribosome | 0.2065 | 1.2 | 0.53856 |
| YDR368W | YPR1 | oxidoreductase | 0.2065 | 1.2 | 0.0905739 |
| YLR216C | CPR6 | unfolded protein binding | 0.2067 | 1.2 | 0.150434 |
| YOR287C | | unknown | 0.2068 | 1.2 | 0.3349 |
| YDR394W | RPT3 | ATPase * | 0.2072 | 1.2 | 0.09225 |
| YKL045W | PR12 | alpha DNA polymerase | 0.2075 | 1.2 | 0.195764 |
| YLL065W | | | 0.2083 | 1.2 | 0.175184 |
| YOR319W | HSH49 | RNA binding | 0.2083 | 1.2 | 0.172456 |
| YMR273C | ZDS1 | protein binding | 0.2083 | 1.2 | 0.128618 |
| YGR072W | UPF3 | unknown | 0.2085 | 1.2 | 0.0940925 |
| YHR142W | CHS7 | unknown | 0.2087 | 1.2 | 0.315467 |
| YMR010W | | unknown | 0.2088 | 1.2 | 0.364446 |
| YER092W | IES5 | unknown | 0.2088 | 1.2 | 0.216276 |
| YOL036W | | unknown | 0.2090 | 1.2 | 0.233145 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YFL052W | | DNA binding | 0.2095 | 1.2 | 0.0119911 |
| YCL059C | KRR1 | unknown | 0.2097 | 1.2 | 0.211965 |
| YDR416W | SYF1 | unknown | 0.2100 | 1.2 | 0.402752 |
| YNL253W | TEX1 | unknown | 0.2107 | 1.2 | 0.49864 |
| YDL053C | PBP4 | unknown | 0.2107 | 1.2 | 0.105205 |
| YDR086C | SSS1 | protein transporter | 0.2110 | 1.2 | 0.235194 |
| YIR024C | | unknown | 0.2110 | 1.2 | 0.0107576 |
| YBR142W | MAK5 | ATP-dependent RNA helicase | 0.2110 | 1.2 | 0.274612 |
| YLR370C | ARC18 | structural constituent of cytoskeleton | 0.2112 | 1.2 | 0.021731 |
| YGL095C | VPS45 | unfolded protein binding | 0.2113 | 1.2 | 0.103886 |
| YLR052W | IES3 | unknown | 0.2115 | 1.2 | 0.55265 |
| YPL011C | TAF3 | general RNA polymerase II transcription factor | 0.2115 | 1.2 | 0.452451 |
| YLR387C | REH1 | unknown | 0.2118 | 1.2 | 0.249962 |
| YMR049C | ERB1 | unknown | 0.2118 | 1.2 | 0.287832 |
| YIL073C | SPO22 | unknown | 0.2118 | 1.2 | 0.288363 |
| YKR074W | | unknown | 0.2120 | 1.2 | 0.573312 |
| YFR027W | ECO1 | acetyltransferase | 0.2122 | 1.2 | 0.47313 |
| YGR079W | | unknown | 0.2122 | 1.2 | 0.624995 |
| YEL074W | | | 0.2123 | 1.2 | 0.00527358 |
| YBR228W | SLX1 | 5'-flap endonuclease | 0.2123 | 1.2 | 0.052611 |
| YOL098C | | metalloendopeptidase | 0.2127 | 1.2 | 0.6106 |
| YKL156W | RPS27A | structural constituent of ribosome | 0.2130 | 1.2 | 0.254447 |
| YEL059C-A | SOM1 | unknown | 0.2130 | 1.2 | 0.230481 |
| YLR166C | SEC10 | protein binding | 0.2130 | 1.2 | 0.486565 |
| YDR078C | SHU2 | unknown | 0.2132 | 1.2 | 0.427389 |
| YDR501W | PLM2 | transcription factor | 0.2132 | 1.2 | 0.456581 |
| YER131W | RPS26B | structural constituent of ribosome | 0.2133 | 1.2 | 0.0268335 |
| YHR039BC | | | 0.2137 | 1.2 | 0.0938629 |
| YKL079W | SMY1 | motor | 0.2137 | 1.2 | 0.362943 |
| YMR203W | TOM40 | protein transporter | 0.2140 | 1.2 | 0.17831 |
| YBR187W | | unknown | 0.2148 | 1.2 | 0.212924 |
| YJR109C | CPA2 | carbamoyl-phosphate synthase (glutamine-hydrolyzing) | 0.2148 | 1.2 | 0.147469 |
| YDR302W | GPI11 | phosphoethanolamine N-methyltransferase | 0.2153 | 1.2 | 0.0635946 |
| YLR276C | DBP9 | ATP-dependent RNA helicase | 0.2160 | 1.2 | 0.106734 |
| YJL138C | TIF2 | translation initiation factor | 0.2162 | 1.2 | 0.363416 |
| YLR442C | SIR3 | histone binding | 0.2162 | 1.2 | 0.600477 |
| YDR447C | RPS17B | structural constituent of ribosome | 0.2165 | 1.2 | 0.047969 |
| YGL185C | | oxidoreductase | 0.2165 | 1.2 | 0.313809 |
| YGR135W | PRE9 | endopeptidase | 0.2173 | 1.2 | 0.0929783 |
| YLR425W | TUS1 | Rho guanyl-nucleotide exchange factor | 0.2175 | 1.2 | 0.144708 |
| YGR153W | | unknown | 0.2175 | 1.2 | 0.473033 |
| YAL041W | CDC24 | signal transducer * | 0.2175 | 1.2 | 0.362508 |
| YJR067C | YAE1 | unknown | 0.2180 | 1.2 | 0.663165 |
| YNL125C | ESBP6 | transporter | 0.2180 | 1.2 | 0.231364 |
| YCL057W | PRD1 | metalloendopeptidase * | 0.2185 | 1.2 | 0.151149 |
| YDR073W | SNF11 | general RNA polymerase II transcription factor | 0.2188 | 1.2 | 0.137427 |
| YML053C | | unknown | 0.2188 | 1.2 | 0.335272 |
| YNL010W | | unknown | 0.2190 | 1.2 | 0.0600224 |
| YGL262W | | unknown | 0.2192 | 1.2 | 0.254889 |
| YCR065W | HCM1 | specific RNA polymerase II transcription factor | 0.2192 | 1.2 | 0.230375 |
| YDR266C | | unknown | 0.2192 | 1.2 | 0.134242 |
| YDL212W | SHR3 | unfolded protein binding | 0.2197 | 1.2 | 0.293455 |
| YER188W | | | 0.2197 | 1.2 | 0.183911 |
| YNL308C | KRI1 | unknown | 0.2197 | 1.2 | 0.052621 |
| YML120C | NDI1 | NADH dehydrogenase | 0.2197 | 1.2 | 0.243432 |
| YAL046C | | unknown | 0.2197 | 1.2 | 0.345238 |
| YHR123W | EPT1 | ethanolaminephosphotransferase | 0.2203 | 1.2 | 0.126889 |
| YJR012C | | unknown | 0.2205 | 1.2 | 0.0856618 |
| YOL094C | RFC4 | DNA clamp loader * | 0.2208 | 1.2 | 0.0917365 |
| YNL079C | TPM1 | actin lateral binding | 0.2210 | 1.2 | 0.182341 |
| YOR213C | SAS5 | acetyltransferase * | 0.2212 | 1.2 | 0.034141 |
| YCR054C | CTR86 | unknown | 0.2215 | 1.2 | 0.138037 |
| YMR312W | ELP6 | RNA polymerase II transcription elongation factor | 0.2218 | 1.2 | 0.561995 |
| YGL091C | NBP35 | ATPase | 0.2220 | 1.2 | 0.409986 |
| YDR065W | | unknown | 0.2222 | 1.2 | 0.593805 |
| YIL052C | RPL34B | structural constituent of ribosome | 0.2223 | 1.2 | 0.133539 |
| YDR067C | OCA6 | unknown | 0.2225 | 1.2 | 0.268057 |
| YPR137W | RRP9 | snoRNA binding | 0.2227 | 1.2 | 0.265159 |
| YDR043C | NRG1 | DNA binding | 0.2228 | 1.2 | 0.141086 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|------------|
| YBR132C | AGP2 | amino acid transporter | 0.2228 | 1.2 | 0.070194 |
| YGR251W | | unknown | 0.2228 | 1.2 | 0.162678 |
| YHR152W | SPO12 | unknown | 0.2232 | 1.2 | 0.0569875 |
| YOL001W | PHO80 | cyclin-dependent protein kinase | 0.2233 | 1.2 | 0.107034 |
| YKL174C | TPO5 | polyamine transporter | 0.2233 | 1.2 | 0.202898 |
| YNL038W | GPI15 | UDP-glycosyltransferase | 0.2237 | 1.2 | 0.40345 |
| YAR031W | PRM9 | unknown | 0.2240 | 1.2 | 0.145567 |
| YDR054C | CDC34 | ubiquitin conjugating enzyme | 0.2242 | 1.2 | 0.00100987 |
| YBR130C | SHE3 | mRNA binding | 0.2243 | 1.2 | 0.196419 |
| YDR165W | TRM82 | protein binding | 0.2253 | 1.2 | 0.474287 |
| YBR264C | YPT10 | GTPase * | 0.2255 | 1.2 | 0.058086 |
| YLR059C | REX2 | 3'-5' exonuclease | 0.2257 | 1.2 | 0.553452 |
| YPR084W | | unknown | 0.2260 | 1.2 | 0.289684 |
| YGL103W | RPL28 | structural constituent of ribosome* | 0.2262 | 1.2 | 0.0353955 |
| YGR188C | BUB1 | protein binding | 0.2262 | 1.2 | 0.168036 |
| YAR015W | ADE1 | phosphoribosylaminoimidazolesuccinocarboxamide synthase | 0.2265 | 1.2 | 0.322197 |
| YOR203W | | | 0.2265 | 1.2 | 0.119555 |
| YDR236C | FMN1 | riboflavin kinase | 0.2268 | 1.2 | 0.530411 |
| YLR100W | ERG27 | 3-keto sterol reductase | 0.2272 | 1.2 | 0.147638 |
| YOL097C | WRS1 | tryptophan-tRNA ligase | 0.2277 | 1.2 | 0.44715 |
| YER094C | PUP3 | endopeptidase | 0.2283 | 1.2 | 0.264687 |
| YMR235C | RNA1 | Ran GTPase activator | 0.2288 | 1.2 | 0.289632 |
| YKL069W | | unknown | 0.2288 | 1.2 | 0.0346682 |
| YNL141W | AAH1 | adenine deaminase | 0.2290 | 1.2 | 0.25245 |
| YCRX12W | | | 0.2300 | 1.2 | 0.249624 |
| YEL075C | | unknown | 0.2305 | 1.2 | 0.0369989 |
| YCR097WB | | | 0.2312 | 1.2 | 0.212512 |
| YDL148C | NOP14 | snoRNA binding | 0.2312 | 1.2 | 0.0426437 |
| YJR056C | | unknown | 0.2317 | 1.2 | 0.373123 |
| YGL148W | ARO2 | chorismate synthase * | 0.2318 | 1.2 | 0.0740175 |
| YNL229C | URE2 | transcription corepressor * | 0.2320 | 1.2 | 0.379889 |
| YJR007W | SUI2 | translation initiation factor | 0.2320 | 1.2 | 0.168411 |
| YDR421W | ARO80 | specific RNA polymerase II transcription factor | 0.2322 | 1.2 | 0.408847 |
| YML056C | IMD4 | IMP dehydrogenase | 0.2328 | 1.2 | 0.253595 |
| YBL095W | | unknown | 0.2332 | 1.2 | 0.0912774 |
| YER051W | JHD1 | histone demethylase (H3-K36 specific) | 0.2332 | 1.2 | 0.31391 |
| Q0060 | AI3 | endonuclease | 0.2335 | 1.2 | 0.00218623 |
| YHR125W | | | 0.2335 | 1.2 | 0.448389 |
| YKL126W | YPK1 | protein serine/threonine kinase | 0.2343 | 1.2 | 0.0489144 |
| YJR144W | MGM101 | DNA binding | 0.2348 | 1.2 | 0.165467 |
| YBR289W | SNF5 | general RNA polymerase II transcription factor | 0.2348 | 1.2 | 0.0054577 |
| YPR028W | YOP1 | protein binding | 0.2352 | 1.2 | 0.0367908 |
| YKR043C | | unknown | 0.2357 | 1.2 | 0.213975 |
| YMR320W | | | 0.2358 | 1.2 | 0.617323 |
| YGL237C | HAP2 | transcriptional activator | 0.2358 | 1.2 | 0.0216083 |
| YGL110C | CUE3 | unknown | 0.2362 | 1.2 | 0.309886 |
| YGL243W | TAD1 | tRNA specific adenosine deaminase | 0.2365 | 1.2 | 0.364074 |
| YOL067C | RTG1 | transcription coactivator | 0.2365 | 1.2 | 0.115311 |
| YJR127C | RSF2 | transcription factor | 0.2365 | 1.2 | 0.0070084 |
| YKL125W | RRN3 | RNA polymerase I transcription factor | 0.2367 | 1.2 | 0.24365 |
| YDR261W-A | | | 0.2372 | 1.2 | 0.196379 |
| YCL050C | APA1 | bis(5'-nucleosyl)-tetraphosphatase | 0.2375 | 1.2 | 0.187713 |
| YLR194C | | structural constituent of cell wall | 0.2375 | 1.2 | 0.00752993 |
| YNL207W | RIO2 | protein kinase | 0.2380 | 1.2 | 0.286799 |
| YPL146C | NOP53 | rRNA binding | 0.2385 | 1.2 | 0.27512 |
| YLR209C | PNP1 | purine-nucleoside phosphorylase | 0.2387 | 1.2 | 0.471536 |
| YOR321W | PMT3 | dolichyl-phosphate-mannose-protein mannosyltransferase | 0.2388 | 1.2 | 0.363515 |
| YDL192W | ARF1 | GTPase | 0.2393 | 1.2 | 0.0132416 |
| YOR369C | RPS12 | structural constituent of ribosome | 0.2397 | 1.2 | 0.70685 |
| YEL025C | | unknown | 0.2398 | 1.2 | 0.0393382 |
| YIL092W | | unknown | 0.2400 | 1.2 | 0.116198 |
| YOR128C | ADE2 | phosphoribosylaminoimidazole carboxylase | 0.2400 | 1.2 | 0.331973 |
| YDL193W | NUS1 | prenyltransferase | 0.2402 | 1.2 | 0.308976 |
| YOR299W | BUD7 | unknown | 0.2413 | 1.2 | 0.0632771 |
| YER146W | LSM5 | RNA binding | 0.2413 | 1.2 | 0.293464 |
| YJL206C-A | | | 0.2417 | 1.2 | 0.473818 |
| YNL001W | DOM34 | unknown | 0.2417 | 1.2 | 0.0676259 |
| YNR010W | CSE2 | RNA polymerase II transcription mediator | 0.2420 | 1.2 | 0.347888 |
| YKL056C | TMA19 | unknown | 0.2425 | 1.2 | 0.00992914 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-------------|
| YEL042W | GDA1 | guanosine-diphosphatase * | 0.2427 | 1.2 | 0.386283 |
| YFR010W | UBP6 | ubiquitin-specific protease | 0.2430 | 1.2 | 0.0131737 |
| YDR425W | SNX41 | protein transporter | 0.2440 | 1.2 | 0.290291 |
| YBR284W | | unknown | 0.2442 | 1.2 | 0.19127 |
| YGL040C | HEM2 | porphobilinogen synthase | 0.2443 | 1.2 | 0.0599176 |
| YAR008W | SEN34 | tRNA-intron endonuclease | 0.2443 | 1.2 | 0.00634203 |
| YGR229C | SMI1 | unknown | 0.2453 | 1.2 | 0.154132 |
| YGR105W | VMA21 | unknown | 0.2457 | 1.2 | 0.13219 |
| YEL007W | | unknown | 0.2462 | 1.2 | 0.0422219 |
| YOR004W | UTP23 | unknown | 0.2462 | 1.2 | 0.480456 |
| YMR217W | GUA1 | GMP synthase (glutamine-hydrolyzing) | 0.2463 | 1.2 | 0.417866 |
| YJL124C | LSM1 | RNA binding | 0.2465 | 1.2 | 0.0799271 |
| YNL166C | BNI5 | unknown | 0.2468 | 1.2 | 0.0145426 |
| YOR194C | TOA1 | general RNA polymerase II transcription factor | 0.2472 | 1.2 | 0.0327548 |
| YMR112C | MED11 | RNA polymerase II transcription mediator | 0.2475 | 1.2 | 0.301758 |
| YFR045W | | transporter | 0.2492 | 1.2 | 0.253317 |
| YCR038C | BUD5 | signal transducer * | 0.2497 | 1.2 | 0.077876 |
| YMR314W | PRE5 | endopeptidase | 0.2502 | 1.2 | 0.127937 |
| YJL183W | MNN11 | alpha-1,6-mannosyltransferase | 0.2503 | 1.2 | 0.158975 |
| YOR340C | RPA43 | DNA-directed RNA polymerase | 0.2508 | 1.2 | 0.376509 |
| YNL118C | DCP2 | mRNA binding | 0.2515 | 1.2 | 0.241346 |
| YBL078C | ATG8 | microtubule binding | 0.2515 | 1.2 | 0.127693 |
| YKL030W | | | 0.2518 | 1.2 | 0.101663 |
| YHR127W | | unknown | 0.2518 | 1.2 | 0.0873626 |
| YER025W | GCD11 | translation initiation factor | 0.2523 | 1.2 | 0.0939522 |
| YKR090W | PXL1 | protein binding | 0.2525 | 1.2 | 0.190128 |
| YOL017W | ESC8 | unknown | 0.2525 | 1.2 | 0.21091 |
| YIL025C | | | 0.2528 | 1.2 | 0.020315 |
| YHR206W | SKN7 | transcription factor | 0.2532 | 1.2 | 0.224089 |
| YBL083C | | | 0.2533 | 1.2 | 0.0765923 |
| YKL193C | SDS22 | enzyme regulator | 0.2535 | 1.2 | 0.11416 |
| YKR012C | | | 0.2545 | 1.2 | 0.418796 |
| YFR016C | | unknown | 0.2548 | 1.2 | 0.132335 |
| YDL105W | NSE4 | unknown | 0.2550 | 1.2 | 0.173526 |
| YDL135C | RD11 | signal transducer * | 0.2552 | 1.2 | 0.0586848 |
| YLR017W | MEU1 | S-methyl-5-thioadenosine phosphorylase | 0.2553 | 1.2 | 0.0523499 |
| YJR001W | AVT1 | neutral amino acid transporter | 0.2565 | 1.2 | 0.0367596 |
| YPL058C | PDR12 | xenobiotic-transporting ATPase * | 0.2565 | 1.2 | 0.189859 |
| YMR240C | CUS1 | protein binding | 0.2567 | 1.2 | 0.133366 |
| YLR118C | | palmitoyl-(protein) hydrolase | 0.2575 | 1.2 | 0.0105155 |
| YLR341W | SPO77 | unknown | 0.2577 | 1.2 | 0.338439 |
| YDR359C | VID21 | unknown | 0.2577 | 1.2 | 0.324662 |
| YML080W | DUS1 | tRNA dihydrouridine synthase | 0.2577 | 1.2 | 0.485112 |
| YBL054W | | unknown | 0.2578 | 1.2 | 0.383595 |
| YCR037C | PHO87 | inorganic phosphate transporter | 0.2578 | 1.2 | 0.300373 |
| YDL106C | PHO2 | transcription factor | 0.2580 | 1.2 | 0.0348297 |
| YDR190C | RVB1 | ATPase * | 0.2583 | 1.2 | 0.144528 |
| YHR048W | | drug transporter | 0.2587 | 1.2 | 0.531096 |
| YER168C | CCA1 | tRNA adenylyltransferase | 0.2588 | 1.2 | 0.0189504 |
| YNL074C | MLF3 | unknown | 0.2592 | 1.2 | 0.158077 |
| YOL040C | RPS15 | structural constituent of ribosome | 0.2593 | 1.2 | 0.0260094 |
| YDR395W | SXM1 | protein carrier | 0.2598 | 1.2 | 0.0992076 |
| YPL017C | | S-adenosylmethionine-dependent methyltransferase | 0.2600 | 1.2 | 0.792581 |
| YDR469W | SDC1 | histone lysine N-methyltransferase (H3-K4 specific) | 0.2602 | 1.2 | 0.00888136 |
| YOR269W | PAC1 | unknown | 0.2602 | 1.2 | 0.314412 |
| YCR087W | | | 0.2603 | 1.2 | 0.0461912 |
| YDR366C | | unknown | 0.2605 | 1.2 | 0.0711693 |
| YKL012W | PRP40 | RNA binding | 0.2607 | 1.2 | 0.165549 |
| YPR203W | | unknown | 0.2608 | 1.2 | 0.162637 |
| YHR090C | YNG2 | enzyme activator | 0.2612 | 1.2 | 0.00951868 |
| YGR096W | TPC1 | transporter | 0.2615 | 1.2 | 0.552549 |
| YPR090W | | | 0.2623 | 1.2 | 0.00706889 |
| YGR276C | RNH70 | 3'-5' exonuclease | 0.2633 | 1.2 | 0.155812 |
| YJL187C | SWE1 | protein kinase | 0.2635 | 1.2 | 0.0525705 |
| YPR063C | | unknown | 0.2642 | 1.2 | 0.0222418 |
| YOR186W | | unknown | 0.2642 | 1.2 | 0.106565 |
| YDL059C | RAD59 | protein binding | 0.2643 | 1.2 | 0.0787789 |
| YNL123W | NMA111 | serine-type peptidase | 0.2643 | 1.2 | 0.158907 |
| YGR106C | | unknown | 0.2643 | 1.2 | 0.000763048 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|------------|
| YBL009W | | protein serine/threonine kinase | 0.2643 | 1.2 | 0.211382 |
| YBR297W | MAL33 | transcription factor | 0.2655 | 1.2 | 0.245612 |
| YPL270W | MDL2 | ATPase , coupled to transmembrane movement of substanc | 0.2655 | 1.2 | 0.134359 |
| YBR292C | | | 0.2655 | 1.2 | 0.364644 |
| YML034W | SRC1 | unknown | 0.2658 | 1.2 | 0.307617 |
| YDR153C | ENT5 | clathrin binding | 0.2668 | 1.2 | 0.0978425 |
| YOR310C | NOP58 | unknown | 0.2683 | 1.2 | 0.173681 |
| YKL181W | PRS1 | ribose phosphate diphosphokinase | 0.2685 | 1.2 | 0.122701 |
| YDR363W | ESC2 | unknown | 0.2687 | 1.2 | 0.00376801 |
| YDR184C | ATC1 | unknown | 0.2690 | 1.2 | 0.11266 |
| YKL155C | RSM22 | structural constituent of ribosome | 0.2695 | 1.2 | 0.112748 |
| YOR343W-A | | | 0.2695 | 1.2 | 0.0992868 |
| YPL199C | | unknown | 0.2697 | 1.2 | 0.0222173 |
| YGL231C | | unknown | 0.2698 | 1.2 | 0.0461556 |
| YFL039C | ACT1 | structural constituent of cytoskeleton | 0.2700 | 1.2 | 0.113005 |
| YMR170C | ALD2 | 3-chloroallyl aldehyde dehydrogenase | 0.2708 | 1.2 | 0.335334 |
| YHR172W | SPC97 | structural constituent of cytoskeleton | 0.2715 | 1.2 | 0.00376801 |
| YDR071C | PAA1 | aralkylamine N-acetyltransferase * | 0.2717 | 1.2 | 0.111615 |
| YDL110C | TMA17 | unknown | 0.2720 | 1.2 | 0.243543 |
| YFL066C | | unknown | 0.2722 | 1.2 | 0.0590227 |
| YDR364C | CDC40 | RNA splicing factor | 0.2725 | 1.2 | 0.240535 |
| YML094W | GIM5 | tubulin binding | 0.2727 | 1.2 | 0.507227 |
| YDL036C | PUS9 | tRNA-pseudouridine synthase | 0.2728 | 1.2 | 0.347938 |
| YMR158W | MRPS8 | structural constituent of ribosome | 0.2728 | 1.2 | 0.0357293 |
| YOL077C | BRX1 | rRNA primary transcript binding | 0.2732 | 1.2 | 0.258736 |
| YDR237W | MRPL7 | structural constituent of ribosome | 0.2733 | 1.2 | 0.191539 |
| YDR282C | | unknown | 0.2738 | 1.2 | 0.183089 |
| YNR023W | SNF12 | general RNA polymerase II transcription factor | 0.2738 | 1.2 | 0.345581 |
| YLR193C | | unknown | 0.2740 | 1.2 | 0.178595 |
| YNL252C | MRPL17 | structural constituent of ribosome | 0.2742 | 1.2 | 0.0310491 |
| YPL258C | THI21 | phosphomethylpyrimidine kinase | 0.2743 | 1.2 | 0.175314 |
| YIL024C | | unknown | 0.2745 | 1.2 | 0.0113285 |
| YCLX10C | | | 0.2747 | 1.2 | 0.362386 |
| YDR324C | UTP4 | snoRNA binding | 0.2747 | 1.2 | 0.175598 |
| YKL209C | STE6 | ATPase , coupled to transmembrane movement of substanc | 0.2748 | 1.2 | 0.198467 |
| YPR162C | ORC4 | DNA replication origin binding | 0.2752 | 1.2 | 0.684754 |
| YDL103C | QRII | UDP-N-acetylglucosamine diphosphorylase | 0.2757 | 1.2 | 0.0108105 |
| YCR009C | RVS161 | cytoskeletal protein binding | 0.2758 | 1.2 | 0.11788 |
| YOL106W | | | 0.2760 | 1.2 | 0.155653 |
| YBR004C | GPI18 | mannosyltransferase | 0.2760 | 1.2 | 0.0305271 |
| YFR048W | RMD8 | unknown | 0.2762 | 1.2 | 0.0340041 |
| YOR163W | DDP1 | diphosphoinositol-polyphosphate diphosphatase | 0.2765 | 1.2 | 0.175881 |
| YDL195W | SEC31 | structural molecule | 0.2767 | 1.2 | 0.195066 |
| YER047C | SAP1 | ATPase | 0.2773 | 1.2 | 0.181525 |
| YMR043W | MCM1 | DNA binding | 0.2777 | 1.2 | 0.235303 |
| YLR383W | SMC6 | unknown | 0.2782 | 1.2 | 0.306265 |
| YDR072C | IPT1 | transferase , transferring phosphorus-containing groups | 0.2782 | 1.2 | 0.252582 |
| YIR009W | MSL1 | RNA binding | 0.2782 | 1.2 | 0.0323755 |
| YOR064C | YNG1 | histone acetyltransferase | 0.2787 | 1.2 | 0.0494563 |
| YDL002C | NHP10 | unknown | 0.2790 | 1.2 | 0.223141 |
| YDL057W | | unknown | 0.2792 | 1.2 | 0.0067468 |
| YDL186W | | unknown | 0.2795 | 1.2 | 0.0292376 |
| YLR248W | RCK2 | protein serine/threonine kinase | 0.2795 | 1.2 | 0.0249379 |
| YOR281C | PLP2 | GTPase inhibitor | 0.2798 | 1.2 | 0.692365 |
| YLR151C | PCD1 | pyrophosphatase | 0.2800 | 1.2 | 0.354762 |
| YJR100C | | unknown | 0.2805 | 1.2 | 0.10521 |
| YLR175W | CBF5 | pseudouridylylate synthase | 0.2807 | 1.2 | 0.161506 |
| YDR238C | SEC26 | unknown | 0.2810 | 1.2 | 0.0429732 |
| YKL080W | VMA5 | hydrogen-transporting ATPase , rotational mechanism | 0.2812 | 1.2 | 0.130258 |
| YGR234W | YHB1 | nitric oxide reductase | 0.2817 | 1.2 | 0.106908 |
| YDR473C | PRP3 | RNA splicing factor | 0.2822 | 1.2 | 0.0489397 |
| YOR316C | COT1 | di-, tri-valent inorganic cation transporter | 0.2825 | 1.2 | 0.0859813 |
| YLR455W | | unknown | 0.2827 | 1.2 | 0.116735 |
| YFL047W | RGD2 | Rho GTPase activator | 0.2828 | 1.2 | 0.0133458 |
| YDL225W | SHS1 | structural constituent of cytoskeleton | 0.2830 | 1.2 | 0.101705 |
| YBR109C | CMD1 | protein binding | 0.2830 | 1.2 | 0.0795603 |
| YGR137W | | | 0.2832 | 1.2 | 0.0829099 |
| YDR163W | CWC15 | unknown | 0.2835 | 1.2 | 0.179771 |
| YMR030W | RSF1 | unknown | 0.2838 | 1.2 | 0.0764922 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|------------|
| YJR085C | | unknown | 0.2842 | 1.2 | 0.063361 |
| YLR229C | CDC42 | GTPase * | 0.2843 | 1.2 | 0.0488772 |
| YGR074W | SMD1 | RNA splicing factor | 0.2845 | 1.2 | 0.28898 |
| YDL108W | KIN28 | general RNA polymerase II transcription factor | 0.2845 | 1.2 | 0.237734 |
| YIL116W | HIS5 | histidinol-phosphate transaminase | 0.2847 | 1.2 | 0.00934731 |
| YJL222W | VTH2 | signal sequence binding | 0.2853 | 1.2 | 0.0252787 |
| YDR045C | RPC11 | DNA-directed RNA polymerase | 0.2853 | 1.2 | 0.041639 |
| YAR019C | CDC15 | protein kinase | 0.2855 | 1.2 | 0.273523 |
| YIL004C | BET1 | v-SNARE | 0.2855 | 1.2 | 0.0786559 |
| YDR468C | TLG1 | v-SNARE * | 0.2857 | 1.2 | 0.0897131 |
| YPL165C | SET6 | unknown | 0.2858 | 1.2 | 0.486303 |
| YBR155W | CNS1 | unfolded protein binding | 0.2858 | 1.2 | 0.0649887 |
| YNL337W | | | 0.2860 | 1.2 | 0.114855 |
| YLR420W | URA4 | dihydroorotase | 0.2860 | 1.2 | 0.0240395 |
| YDR058C | TGL2 | triacylglycerol lipase | 0.2868 | 1.2 | 0.122623 |
| YIL096C | | unknown | 0.2872 | 1.2 | 0.215658 |
| YBR071W | | unknown | 0.2875 | 1.2 | 0.290717 |
| YDL068W | | | 0.2878 | 1.2 | 0.149483 |
| YDR097C | MSH6 | DNA binding | 0.2897 | 1.2 | 0.0608023 |
| YLR104W | | unknown | 0.2898 | 1.2 | 0.0816736 |
| YNL260C | | unknown | 0.2902 | 1.2 | 0.151976 |
| YEL019C | MMS21 | SUMO ligase | 0.2908 | 1.2 | 0.0563927 |
| YER085C | | unknown | 0.2913 | 1.2 | 0.089874 |
| YCR007C | | unknown | 0.2918 | 1.2 | 0.196194 |
| YER071C | | unknown | 0.2927 | 1.2 | 0.0893787 |
| YNL002C | RLP7 | rRNA binding | 0.2930 | 1.2 | 0.181953 |
| YOL003C | PFA4 | protein-cysteine S-palmitoyltransferase | 0.2930 | 1.2 | 0.0812157 |
| YGL179C | TOS3 | protein kinase | 0.2932 | 1.2 | 0.138669 |
| YHR162W | | unknown | 0.2933 | 1.2 | 0.0851827 |
| YNL065W | AQR1 | monocarboxylic acid transporter | 0.2933 | 1.2 | 0.212312 |
| YLR432W | IMD3 | IMP dehydrogenase | 0.2935 | 1.2 | 0.290719 |
| YFR006W | | X-Pro aminopeptidase | 0.2937 | 1.2 | 0.0226186 |
| YBL040C | ERD2 | HDEL sequence binding | 0.2938 | 1.2 | 0.0312894 |
| YDR543C | | | 0.2942 | 1.2 | 0.233374 |
| YJL122W | ALB1 | unknown | 0.2945 | 1.2 | 0.1925 |
| YMR176W | ECM5 | unknown | 0.2947 | 1.2 | 0.144141 |
| YDL201W | TRM8 | protein binding | 0.2948 | 1.2 | 0.302221 |
| YML101C | CUE4 | unknown | 0.2953 | 1.2 | 0.0523719 |
| YOR311C | HSD1 | unknown | 0.2958 | 1.2 | 0.12189 |
| YJL005W | CYR1 | adenylate cyclase | 0.2960 | 1.2 | 0.180368 |
| YDR365W-A | | | 0.2968 | 1.2 | 0.188724 |
| YER039C | HVG1 | unknown | 0.2973 | 1.2 | 0.182631 |
| YNL147W | LSM7 | RNA binding | 0.2973 | 1.2 | 0.0275838 |
| YHR170W | NMD3 | protein binding | 0.2973 | 1.2 | 0.191785 |
| YJR082C | EAF6 | unknown | 0.2975 | 1.2 | 0.213894 |
| YGR252W | GCN5 | histone acetyltransferase * | 0.2985 | 1.2 | 0.480719 |
| YNL083W | SAL1 | transporter | 0.2988 | 1.2 | 0.217947 |
| YGR028W | MSP1 | ATPase | 0.3008 | 1.2 | 0.150702 |
| YDL165W | CDC36 | 3'-5'-exoribonuclease | 0.3010 | 1.2 | 0.333445 |
| YCR042C | TAF2 | general RNA polymerase II transcription factor | 0.3013 | 1.2 | 0.0534277 |
| YOL042W | NGL1 | endonuclease | 0.3020 | 1.2 | 0.060534 |
| YMR263W | SAP30 | histone deacetylase | 0.3025 | 1.2 | 0.0207154 |
| YDL213C | NOP6 | unknown | 0.3037 | 1.2 | 0.041178 |
| YDL217C | TIM22 | protein transporter | 0.3040 | 1.2 | 0.511644 |
| YBR082C | UBC4 | ubiquitin conjugating enzyme | 0.3050 | 1.2 | 0.0192093 |
| YBR156C | SLI15 | protein kinase | 0.3058 | 1.2 | 0.0397654 |
| YDL226C | GCS1 | actin binding | 0.3060 | 1.2 | 0.00732041 |
| YLR293C | GSP1 | GTPase | 0.3062 | 1.2 | 0.0663701 |
| YHL031C | GOS1 | v-SNARE | 0.3065 | 1.2 | 0.10777 |
| YMR288W | HSH155 | mRNA binding | 0.3067 | 1.2 | 0.456591 |
| YLR227C | ADY4 | structural molecule | 0.3067 | 1.2 | 0.0881128 |
| YNL183C | NPR1 | kinase | 0.3070 | 1.2 | 0.382339 |
| YBR137W | | unknown | 0.3078 | 1.2 | 0.0516986 |
| YIL027C | KRE27 | unknown | 0.3083 | 1.2 | 0.0359658 |
| YKR061W | KTR2 | mannosyltransferase | 0.3083 | 1.2 | 0.0162137 |
| YDR056C | | unknown | 0.3088 | 1.2 | 0.0200136 |
| YMR079W | SEC14 | phosphatidylinositol transporter | 0.3095 | 1.2 | 0.100294 |
| YPL060W | LPE10 | magnesium ion transporter | 0.3095 | 1.2 | 0.0244191 |
| YDR084C | TVP23 | unknown | 0.3095 | 1.2 | 0.0297584 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|---|----------------------------|-------------|------------|
| YAR023C | | unknown | 0.3095 | 1.2 | 0.137216 |
| YMR275C | BUL1 | protein binding | 0.3097 | 1.2 | 0.140026 |
| YBR257W | POP4 | ribonuclease P * | 0.3098 | 1.2 | 0.0131032 |
| YLR180W | SAM1 | methionine adenosyltransferase | 0.3102 | 1.2 | 0.146012 |
| YER006W | NUG1 | GTPase | 0.3103 | 1.2 | 0.151331 |
| YGL116W | CDC20 | enzyme activator | 0.3107 | 1.2 | 0.10693 |
| YHR200W | RPN10 | endopeptidase | 0.3108 | 1.2 | 0.0186369 |
| YAL015C | NTG1 | DNA-(apurinic or apyrimidinic site) lyase * | 0.3112 | 1.2 | 0.00146686 |
| YBL076C | ILS1 | isoleucine-tRNA ligase | 0.3112 | 1.2 | 0.188992 |
| YLR309C | IMH1 | unknown | 0.3118 | 1.2 | 0.228186 |
| YDR189W | SLY1 | SNARE binding | 0.3120 | 1.2 | 0.358734 |
| YBR246W | | unknown | 0.3135 | 1.2 | 0.0506469 |
| YLR380W | CSR1 | phosphatidylinositol transporter | 0.3137 | 1.2 | 0.219645 |
| YMR259C | | unknown | 0.3142 | 1.2 | 0.0811376 |
| YFL031W | HAC1 | transcription factor | 0.3143 | 1.2 | 0.15147 |
| YBR011C | IPP1 | inorganic diphosphatase | 0.3147 | 1.2 | 0.0847703 |
| YBR203W | COS111 | unknown | 0.3148 | 1.2 | 0.0659107 |
| YOR239W(1) | | | 0.3152 | 1.2 | 0.436596 |
| YOR244W | ESA1 | histone acetyltransferase | 0.3153 | 1.2 | 0.146922 |
| YFL025C | BST1 | carboxylic ester hydrolase | 0.3158 | 1.2 | 0.298488 |
| YDL130W | RPP1B | structural constituent of ribosome | 0.3168 | 1.2 | 0.120698 |
| YER088C | DOT6 | unknown | 0.3172 | 1.2 | 0.0456176 |
| YBL022C | PIM1 | ATP-dependent peptidase | 0.3172 | 1.2 | 0.049749 |
| YNL313C | | unknown | 0.3183 | 1.2 | 0.554478 |
| YML016C | PPZ1 | protein serine/threonine phosphatase | 0.3187 | 1.2 | 0.0339913 |
| YIL103W | DPH1 | unknown | 0.3190 | 1.2 | 0.0440198 |
| YGL126W | SCS3 | unknown | 0.3192 | 1.2 | 0.186075 |
| YHL021C | FMP12 | unknown | 0.3195 | 1.2 | 0.0944435 |
| YLR243W | | signal sequence binding | 0.3197 | 1.2 | 0.0187408 |
| YNR027W | BUD17 | unknown | 0.3198 | 1.2 | 0.209163 |
| YMR309C | NIP1 | translation initiation factor | 0.3200 | 1.2 | 0.106909 |
| YOR048C | RAT1 | 5'-3' exoribonuclease | 0.3203 | 1.2 | 0.00405686 |
| YBR165W | UBS1 | unknown | 0.3203 | 1.2 | 0.00691753 |
| YIL028W | | | 0.3205 | 1.2 | 0.181484 |
| YKL119C | VPH2 | unknown | 0.3207 | 1.2 | 0.464783 |
| YLR368W | MDM30 | unknown | 0.3208 | 1.2 | 0.234646 |
| YJR052W | RAD7 | DNA binding | 0.3210 | 1.2 | 0.0507301 |
| YAR066W | | unknown | 0.3220 | 1.3 | 0.277759 |
| YOL090W | MSH2 | ATPase * | 0.3228 | 1.3 | 0.173821 |
| YPL207W | TYW1 | unknown | 0.3235 | 1.3 | 0.368234 |
| YCL066W | HMLALPHA1 | transcription coactivator | 0.3235 | 1.3 | 0.444371 |
| YER012W | PRE1 | endopeptidase | 0.3240 | 1.3 | 0.0115158 |
| YHR154W | RTT107 | unknown | 0.3242 | 1.3 | 0.0250395 |
| YJL006C | CTK2 | cyclin-dependent protein kinase | 0.3242 | 1.3 | 0.465752 |
| YDR034C-C | | | 0.3248 | 1.3 | 0.00148546 |
| YLR291C | GCD7 | translation initiation factor | 0.3252 | 1.3 | 0.213169 |
| YNR075W | COS10 | unknown | 0.3255 | 1.3 | 0.105745 |
| YIL138C | TPM2 | actin lateral binding | 0.3257 | 1.3 | 0.153284 |
| YDR047W | HEM12 | uroporphyrinogen decarboxylase | 0.3257 | 1.3 | 0.0167387 |
| YEL024W | RIP1 | ubiquinol-cytochrome-c reductase | 0.3267 | 1.3 | 0.0137258 |
| YGL060W | YBP2 | unknown | 0.3268 | 1.3 | 0.0483485 |
| YNL073W | MSK1 | lysine-tRNA ligase | 0.3270 | 1.3 | 0.0441222 |
| YOL061W | PRS5 | ribose phosphate diphosphokinase | 0.3275 | 1.3 | 0.00448467 |
| YLR303W | MET17 | cysteine synthase * | 0.3277 | 1.3 | 0.0929698 |
| YMR210W | | unknown | 0.3282 | 1.3 | 0.244544 |
| YJL112W | MDV1 | unknown | 0.3285 | 1.3 | 0.00071224 |
| YMR267W | PPA2 | inorganic diphosphatase | 0.3290 | 1.3 | 0.104133 |
| YNL263C | YIF1 | unknown | 0.3290 | 1.3 | 0.00195159 |
| YBL045C | COR1 | ubiquinol-cytochrome-c reductase | 0.3298 | 1.3 | 0.0531117 |
| YNL284C-A | | | 0.3303 | 1.3 | 0.113605 |
| YKL064W | MNR2 | magnesium ion transporter | 0.3305 | 1.3 | 0.0660159 |
| YNL106C | INP52 | inositol-polyphosphate 5-phosphatase | 0.3305 | 1.3 | 0.264571 |
| YMR215W | GAS3 | 1,3-beta-glucanosyltransferase | 0.3305 | 1.3 | 0.15332 |
| YOL127W | RPL25 | structural constituent of ribosome* | 0.3308 | 1.3 | 0.0984725 |
| YGR057C | LST7 | protein transporter | 0.3315 | 1.3 | 0.249987 |
| YNL154C | YCK2 | casein kinase | 0.3317 | 1.3 | 0.0456629 |
| YLR019W | PSR2 | phosphoprotein phosphatase | 0.3317 | 1.3 | 0.0234776 |
| YPR165W | RHO1 | GTPase * | 0.3317 | 1.3 | 0.163847 |
| YBR199W | KTR4 | mannosyltransferase | 0.3323 | 1.3 | 0.162804 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YML011C | RAD33 | unknown | 0.3330 | 1.3 | 0.282426 |
| YCL024W | KCC4 | protein kinase | 0.3337 | 1.3 | 0.0558064 |
| YDR502C | SAM2 | methionine adenosyltransferase | 0.3337 | 1.3 | 0.0322586 |
| YLR195C | NMT1 | glycylpeptide N-tetradecanoyltransferase | 0.3340 | 1.3 | 0.0724227 |
| YCL037C | SRO9 | RNA binding | 0.3342 | 1.3 | 0.211033 |
| YIL020C | HIS6 | 1-(5-phosphoribosyl)-5-[(5-phosphoribosylamino)methylid | 0.3343 | 1.3 | 0.0961617 |
| YOR276W | CAF20 | translation regulator | 0.3348 | 1.3 | 0.0793003 |
| YLR256W-A | | | 0.3350 | 1.3 | 0.152657 |
| YGR245C | SDA1 | unknown | 0.3357 | 1.3 | 0.509982 |
| YPL194W | DDC1 | unknown | 0.3360 | 1.3 | 0.446199 |
| YBR034C | HMT1 | protein-arginine N-methyltransferase | 0.3372 | 1.3 | 0.0100556 |
| YPL009C | | unknown | 0.3372 | 1.3 | 0.285743 |
| YGR277C | | panetheine-phosphate adenyltransferase | 0.3377 | 1.3 | 9.26074E-05 |
| YBR293W | VBA2 | multidrug transporter | 0.3377 | 1.3 | 0.102285 |
| YNL054W-A | | | 0.3395 | 1.3 | 0.0759197 |
| YNR015W | SMM1 | tRNA dihydrouridine synthase | 0.3397 | 1.3 | 0.487621 |
| YKL099C | UTP11 | snoRNA binding | 0.3400 | 1.3 | 0.229313 |
| YDR336W | | unknown | 0.3407 | 1.3 | 0.269227 |
| YDR064W | RPS13 | structural constituent of ribosome | 0.3408 | 1.3 | 0.0417386 |
| YJL161W | FMP33 | unknown | 0.3412 | 1.3 | 0.160599 |
| YNR074C | AIF1 | oxidoreductase | 0.3412 | 1.3 | 0.152855 |
| YNL050C | | unknown | 0.3413 | 1.3 | 0.0712413 |
| YDR083W | RRP8 | methyltransferase | 0.3417 | 1.3 | 0.153002 |
| YPR045C | | unknown | 0.3420 | 1.3 | 0.327041 |
| YJL110C | GZF3 | transcription factor | 0.3422 | 1.3 | 0.245413 |
| YDL198C | GGC1 | guanine nucleotide transporter | 0.3425 | 1.3 | 0.0371653 |
| YNL142W | MEP2 | ammonium transporter | 0.3428 | 1.3 | 0.0610554 |
| YNL110C | NOP15 | unknown | 0.3430 | 1.3 | 0.229352 |
| YNL180C | RHO5 | GTPase | 0.3440 | 1.3 | 0.0248899 |
| YMR178W | | unknown | 0.3442 | 1.3 | 0.0126728 |
| YDL028C | MPS1 | protein threonine/tyrosine kinase | 0.3460 | 1.3 | 0.0722155 |
| YOR047C | STD1 | protein kinase | 0.3462 | 1.3 | 0.301447 |
| YDL045C | FAD1 | FMN adenyltransferase | 0.3463 | 1.3 | 0.0154089 |
| YLR074C | BUD20 | unknown | 0.3465 | 1.3 | 0.0329259 |
| YKR025W | RPC37 | DNA-directed RNA polymerase | 0.3472 | 1.3 | 0.0586111 |
| YNL181W | | oxidoreductase | 0.3473 | 1.3 | 0.0774725 |
| YMR298W | LIP1 | sphingosine N-acyltransferase | 0.3475 | 1.3 | 0.0671655 |
| YER137C | | unknown | 0.3478 | 1.3 | 0.134993 |
| YOR007C | SGT2 | unknown | 0.3478 | 1.3 | 0.112838 |
| YAR028W | | unknown | 0.3480 | 1.3 | 0.110732 |
| YLR038C | COX12 | cytochrome-c oxidase | 0.3495 | 1.3 | 0.375377 |
| YML097C | VPS9 | guanyl-nucleotide exchange factor | 0.3510 | 1.3 | 0.100685 |
| YHR211W | FLO5 | mannose binding | 0.3510 | 1.3 | 0.20237 |
| YLR060W | FRS1 | phenylalanine-tRNA ligase | 0.3512 | 1.3 | 0.0252629 |
| YBR196C | PGI1 | glucose-6-phosphate isomerase | 0.3520 | 1.3 | 0.169196 |
| YFL023W | BUD27 | unknown | 0.3523 | 1.3 | 0.125296 |
| YPR109W | | unknown | 0.3527 | 1.3 | 0.0115682 |
| YGL115W | SNF4 | protein kinase | 0.3530 | 1.3 | 0.0479353 |
| YHR091C | MSR1 | arginine-tRNA ligase | 0.3535 | 1.3 | 0.277625 |
| YNL251C | NRD1 | RNA binding | 0.3537 | 1.3 | 0.294214 |
| YLR335W | NUP2 | structural molecule | 0.3545 | 1.3 | 0.212291 |
| YNL220W | ADE12 | adenylosuccinate synthase | 0.3545 | 1.3 | 0.153954 |
| YDR089W | | unknown | 0.3547 | 1.3 | 0.265498 |
| YGR138C | TPO2 | spermine transporter | 0.3552 | 1.3 | 0.00530685 |
| YNL309W | STB1 | transcriptional activator | 0.3553 | 1.3 | 0.326359 |
| YJL118W | | unknown | 0.3553 | 1.3 | 0.171763 |
| YGR058W | | unknown | 0.3558 | 1.3 | 0.181036 |
| YFR004W | RPN11 | endopeptidase | 0.3578 | 1.3 | 0.00633033 |
| YER027C | GAL83 | AMP-activated protein kinase | 0.3582 | 1.3 | 0.0759152 |
| YMR005W | TAF4 | RNA polymerase II transcription factor | 0.3585 | 1.3 | 0.0781685 |
| YKL088W | | purine nucleotide binding | 0.3585 | 1.3 | 0.164433 |
| YOL144W | NOP8 | unknown | 0.3588 | 1.3 | 0.465894 |
| YHR106W | TRR2 | thioredoxin-disulfide reductase | 0.3590 | 1.3 | 0.0814045 |
| YOL043C | NTG2 | DNA-(apurinic or apyrimidinic site) lyase * | 0.3592 | 1.3 | 0.127536 |
| YGL029W | CGR1 | unknown | 0.3595 | 1.3 | 0.0571443 |
| YDL227C | HO | endonuclease | 0.3598 | 1.3 | 0.0630797 |
| YHR061C | GIC1 | small GTPase regulator | 0.3607 | 1.3 | 0.103102 |
| YLR064W | | unknown | 0.3610 | 1.3 | 0.0182285 |
| YDL230W | PTP1 | protein tyrosine phosphatase | 0.3610 | 1.3 | 0.144822 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|------------|
| YML012W | ERV25 | unknown | 0.3610 | 1.3 | 0.160971 |
| YFL053W | DAK2 | glycerone kinase | 0.3615 | 1.3 | 0.143986 |
| YER032W | FIR1 | unknown | 0.3620 | 1.3 | 0.199377 |
| YPL078C | ATP4 | structural molecule * | 0.3627 | 1.3 | 0.0119898 |
| YDR375C | BCS1 | ATPase | 0.3633 | 1.3 | 0.137506 |
| YJL051W | IRC8 | unknown | 0.3638 | 1.3 | 0.0194293 |
| YDR513W | GRX2 | thiol-disulfide exchange intermediate * | 0.3642 | 1.3 | 0.10078 |
| YLR197W | SIK1 | unknown | 0.3643 | 1.3 | 0.125653 |
| YOR231W | MKK1 | MAP kinase | 0.3648 | 1.3 | 0.180798 |
| YGL073W | HSF1 | transcription factor | 0.3648 | 1.3 | 0.0786332 |
| YOL089C | HAL9 | specific RNA polymerase II transcription factor | 0.3650 | 1.3 | 0.00180215 |
| YER086W | ILV1 | threonine ammonia-lyase | 0.3653 | 1.3 | 0.0356812 |
| YDL122W | UBP1 | ubiquitin-specific protease | 0.3658 | 1.3 | 0.142034 |
| YHR065C | RRP3 | ATP-dependent RNA helicase | 0.3667 | 1.3 | 0.0686174 |
| YDR429C | TIF35 | translation initiation factor | 0.3668 | 1.3 | 0.107343 |
| YJR010C-A | SPC1 | unknown | 0.3668 | 1.3 | 0.00503789 |
| YFL040W | | unknown | 0.3670 | 1.3 | 0.10378 |
| YLR264W | RPS28B | structural constituent of ribosome | 0.3678 | 1.3 | 0.173261 |
| YGR253C | PUP2 | endopeptidase | 0.3682 | 1.3 | 0.0531721 |
| YDR392W | SPT3 | transcription cofactor | 0.3685 | 1.3 | 0.00165205 |
| YJR152W | DAL5 | allantoate transporter | 0.3690 | 1.3 | 0.111736 |
| YLR226W | BUR2 | cyclin-dependent protein kinase | 0.3697 | 1.3 | 0.0449872 |
| YNL227C | JJJ1 | unknown | 0.3698 | 1.3 | 0.195103 |
| YFR033C | QCR6 | ubiquinol-cytochrome-c reductase | 0.3702 | 1.3 | 0.0543097 |
| YDR314C | RAD34 | unknown | 0.3707 | 1.3 | 0.0437755 |
| YBR061C | TRM7 | tRNA methyltransferase | 0.3713 | 1.3 | 0.0909942 |
| YDL123W | SNA4 | unknown | 0.3715 | 1.3 | 0.177047 |
| YDL070W | BDF2 | unknown | 0.3717 | 1.3 | 0.0771189 |
| YKL208W | CBT1 | unknown | 0.3720 | 1.3 | 0.271215 |
| YGR037C | ACB1 | long-chain fatty acid transporter | 0.3722 | 1.3 | 0.148607 |
| YDR001C | NTH1 | alpha,alpha-trehalase | 0.3725 | 1.3 | 0.0461789 |
| YFL022C | FRS2 | phenylalanine-tRNA ligase | 0.3730 | 1.3 | 0.0541346 |
| YBR194W | SOY1 | unknown | 0.3735 | 1.3 | 0.0361311 |
| YDR201W | SPC19 | structural constituent of cytoskeleton | 0.3735 | 1.3 | 0.0611205 |
| YPL104W | MSD1 | aspartate-tRNA ligase | 0.3742 | 1.3 | 0.0446667 |
| YBR062C | | unknown | 0.3745 | 1.3 | 0.0511 |
| YEL051W | VMA8 | hydrogen-transporting ATPase , rotational mechanism | 0.3747 | 1.3 | 0.0221746 |
| YDR329C | PEX3 | protein binding | 0.3748 | 1.3 | 0.00340154 |
| YFR034C | PHO4 | transcription factor | 0.3753 | 1.3 | 0.405673 |
| YCR002C | CDC10 | GTPase * | 0.3763 | 1.3 | 0.0390753 |
| YDR430C | CYM1 | metalloendopeptidase | 0.3763 | 1.3 | 0.244077 |
| YPR085C | | unknown | 0.3775 | 1.3 | 0.0289033 |
| YGL155W | CDC43 | signal transducer * | 0.3778 | 1.3 | 0.109475 |
| YCR083W | TRX3 | thiol-disulfide exchange intermediate | 0.3778 | 1.3 | 0.117739 |
| YLR268W | SEC22 | v-SNARE | 0.3785 | 1.3 | 0.0115942 |
| YDR350C | TCM10 | unknown | 0.3787 | 1.3 | 0.103499 |
| YLR265C | NEJ1 | unknown | 0.3797 | 1.3 | 0.0452802 |
| YDR272W | GLO2 | hydroxyacylglutathione hydrolase | 0.3807 | 1.3 | 0.0121378 |
| YNL284C | MRPL10 | structural constituent of ribosome | 0.3807 | 1.3 | 0.0146558 |
| YJR019C | TES1 | acyl-CoA thioesterase | 0.3807 | 1.3 | 0.0481517 |
| YHR110W | ERP5 | unknown | 0.3808 | 1.3 | 0.0633788 |
| YNR034W | SOL1 | unknown | 0.3815 | 1.3 | 0.0465103 |
| YKR080W | MTD1 | methylenetetrahydrofolate dehydrogenase | 0.3815 | 1.3 | 0.159451 |
| YPL052W | OAZ1 | enzyme inhibitor | 0.3822 | 1.3 | 0.0488677 |
| YEL072W | RMD6 | unknown | 0.3823 | 1.3 | 0.126648 |
| YOR237W | HES1 | oxysterol binding | 0.3825 | 1.3 | 0.260569 |
| YDR112W | | | 0.3827 | 1.3 | 0.0775612 |
| YKL081W | TEF4 | translation elongation factor | 0.3828 | 1.3 | 0.00256069 |
| YDL020C | RPN4 | transcriptional activator | 0.3832 | 1.3 | 0.0735858 |
| YBR087W | RFC5 | DNA clamp loader | 0.3832 | 1.3 | 0.00212711 |
| YDR389W | SAC7 | signal transducer * | 0.3833 | 1.3 | 0.0822193 |
| YAR033W | MST28 | protein binding | 0.3840 | 1.3 | 0.151429 |
| YJR097W | JJJ3 | unknown | 0.3842 | 1.3 | 0.0780038 |
| YER133W | GLC7 | protein phosphatase type 1 | 0.3845 | 1.3 | 0.00582644 |
| YJR022W | LSM8 | RNA binding | 0.3857 | 1.3 | 0.454422 |
| YNR028W | CPR8 | peptidyl-prolyl cis-trans isomerase | 0.3860 | 1.3 | 0.0554568 |
| YJR162C | | | 0.3865 | 1.3 | 0.360675 |
| YBR210W | ERV15 | unknown | 0.3870 | 1.3 | 0.0141565 |
| YHR209W | | S-adenosylmethionine-dependent methyltransferase | 0.3877 | 1.3 | 0.0126853 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-------------|
| YGR233C | PHO81 | cyclin-dependent protein kinase | 0.3878 | 1.3 | 0.241326 |
| YPL168W | | unknown | 0.3892 | 1.3 | 0.0693092 |
| YJR113C | RSM7 | structural constituent of ribosome | 0.3892 | 1.3 | 0.0766389 |
| YDR527W | RBA50 | DNA-directed RNA polymerase | 0.3895 | 1.3 | 0.0481616 |
| YMR022W | QRI8 | ubiquitin conjugating enzyme | 0.3897 | 1.3 | 0.00130401 |
| YJL133W | MRS3 | iron ion transporter | 0.3898 | 1.3 | 0.0345282 |
| YJR078W | BNA2 | tryptophan 2,3-dioxygenase | 0.3903 | 1.3 | 0.3433 |
| YOL143C | RIB4 | 6,7-dimethyl-8-ribityllumazine synthase | 0.3908 | 1.3 | 0.0219818 |
| YFL061W | DDI2 | unknown | 0.3908 | 1.3 | 0.00674054 |
| YHL024W | RIM4 | RNA binding | 0.3922 | 1.3 | 0.259553 |
| YGL099W | LSG1 | GTPase | 0.3923 | 1.3 | 0.387602 |
| YBR055C | PRP6 | RNA splicing factor | 0.3933 | 1.3 | 0.0939433 |
| YAL003W | EFB1 | translation elongation factor | 0.3933 | 1.3 | 0.00260375 |
| YDL136W | RPL35B | structural constituent of ribosome | 0.3935 | 1.3 | 0.107267 |
| YNL315C | ATP11 | unfolded protein binding | 0.3950 | 1.3 | 0.148398 |
| YAL033W | POP5 | ribonuclease P * | 0.3952 | 1.3 | 0.108664 |
| YDR145W | TAF12 | general RNA polymerase II transcription factor | 0.3953 | 1.3 | 0.0462536 |
| YBR012C | | | 0.3958 | 1.3 | 0.13285 |
| YGL064C | MRH4 | RNA helicase | 0.3962 | 1.3 | 0.229261 |
| YGL011C | SCL1 | endopeptidase | 0.3972 | 1.3 | 0.03885 |
| YDL161W | ENT1 | clathrin binding | 0.3975 | 1.3 | 0.0950582 |
| YFR008W | FAR7 | unknown | 0.3978 | 1.3 | 0.0522157 |
| YLR328W | NMA1 | nicotinamide-nucleotide adenylyltransferase | 0.3980 | 1.3 | 0.000314098 |
| YDL205C | HEM3 | hydroxymethylbilane synthase | 0.3985 | 1.3 | 0.18973 |
| YAR068W | | unknown | 0.3987 | 1.3 | 0.0896942 |
| YNL322C | KRE1 | structural constituent of cell wall | 0.3988 | 1.3 | 0.0454274 |
| YOR085W | OST3 | dolichyl-diphosphooligosaccharide-protein glycotransferase | 0.3992 | 1.3 | 0.0604623 |
| YPR057W | BRR1 | RNA binding | 0.3997 | 1.3 | 0.35368 |
| YDR440W | DOT1 | protein-lysine N-methyltransferase | 0.3997 | 1.3 | 0.127816 |
| YPL225W | | unknown | 0.3997 | 1.3 | 0.0290287 |
| YBR291C | CTP1 | tricarboxylate carrier | 0.3998 | 1.3 | 0.0126561 |
| YNL244C | SUI1 | translation initiation factor | 0.4002 | 1.3 | 0.0456726 |
| YOL053W | | unknown | 0.4003 | 1.3 | 0.151764 |
| YPL238C | | | 0.4007 | 1.3 | 0.0613746 |
| YPL252C | YAH1 | electron carrier | 0.4013 | 1.3 | 0.209866 |
| YDR337W | MRPS28 | structural constituent of ribosome* | 0.4013 | 1.3 | 0.0752444 |
| YMR257C | PET111 | translation regulator | 0.4015 | 1.3 | 0.202786 |
| YDR280W | RRP45 | 3'-5'-exoribonuclease | 0.4022 | 1.3 | 0.168241 |
| YOR309C | | | 0.4028 | 1.3 | 0.0527659 |
| YDL209C | CWC2 | unknown | 0.4028 | 1.3 | 0.0628444 |
| YDR139C | RUB1 | protein tag | 0.4033 | 1.3 | 0.021619 |
| YDR351W | SBE2 | unknown | 0.4045 | 1.3 | 0.038492 |
| YDL158C | | | 0.4045 | 1.3 | 0.0276075 |
| YKL114C | APN1 | DNA-(apurinic or apyrimidinic site) lyase | 0.4048 | 1.3 | 0.238965 |
| YEL073C | | unknown | 0.4048 | 1.3 | 0.0302557 |
| YHR143W-A | RPC10 | DNA-directed RNA polymerase | 0.4057 | 1.3 | 0.0682927 |
| YJL189W | RPL39 | structural constituent of ribosome | 0.4060 | 1.3 | 0.0919081 |
| YIL034C | CAP2 | actin filament binding | 0.4060 | 1.3 | 0.0395035 |
| YDR261C-C | | | 0.4065 | 1.3 | 0.128247 |
| YOR147W | MDM32 | unknown | 0.4070 | 1.3 | 0.0430178 |
| YDR253C | MET32 | DNA binding | 0.4075 | 1.3 | 0.0526674 |
| YKR052C | MRS4 | iron ion transporter | 0.4082 | 1.3 | 0.0952343 |
| YDR459C | PFA5 | protein-cysteine S-palmitoyltransferase | 0.4090 | 1.3 | 0.210381 |
| YPL129W | TAF14 | general RNA polymerase II transcription factor | 0.4093 | 1.3 | 0.262608 |
| YHR187W | IKI1 | RNA polymerase II transcription elongation factor | 0.4097 | 1.3 | 0.0133826 |
| YBR290W | BSD2 | unknown | 0.4115 | 1.3 | 0.0207699 |
| YAL039C | CYC3 | holocytochrome-c synthase | 0.4118 | 1.3 | 0.000928542 |
| YDR397C | NCB2 | transcription corepressor | 0.4133 | 1.3 | 0.0145439 |
| YBR252W | DUT1 | dUTP diphosphatase | 0.4142 | 1.3 | 0.0325705 |
| YDR441C | APT2 | unknown | 0.4143 | 1.3 | 0.12869 |
| YOR192C-A | | | 0.4147 | 1.3 | 0.0371685 |
| YOL102C | TPT1 | tRNA 2'-phosphotransferase | 0.4150 | 1.3 | 0.318881 |
| YBR033W | EDS1 | unknown | 0.4152 | 1.3 | 0.0171443 |
| YDL079C | MRK1 | glycogen synthase kinase | 0.4157 | 1.3 | 0.0267702 |
| YBR186W | PCH2 | unknown | 0.4158 | 1.3 | 0.125427 |
| YJL011C | RPC17 | DNA-directed RNA polymerase | 0.4158 | 1.3 | 0.0575168 |
| YGR182C | | | 0.4162 | 1.3 | 0.220367 |
| YBL093C | ROX3 | RNA polymerase II transcription mediator | 0.4167 | 1.3 | 0.311958 |
| YNR026C | SEC12 | guanyl-nucleotide exchange factor | 0.4173 | 1.3 | 0.0733029 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-------------|
| YDL018C | ERP3 | unknown | 0.4175 | 1.3 | 0.0775042 |
| YNL175C | NOP13 | RNA binding | 0.4177 | 1.3 | 0.023322 |
| YNL075W | IMP4 | rRNA primary transcript binding | 0.4180 | 1.3 | 0.0497298 |
| YPL024W | RMI1 | unknown | 0.4183 | 1.3 | 0.151392 |
| YNL161W | CBK1 | protein kinase | 0.4185 | 1.3 | 0.0107181 |
| YAR075W | | | 0.4190 | 1.3 | 0.148769 |
| YDR524C | AGE1 | ARF GTPase activator | 0.4192 | 1.3 | 0.109806 |
| YDR405W | MRP20 | structural constituent of ribosome | 0.4195 | 1.3 | 0.00542521 |
| YKL063C | | unknown | 0.4197 | 1.3 | 0.10551 |
| YDR470C | UGO1 | transporter | 0.4203 | 1.3 | 0.0573441 |
| YOR262W | | unknown | 0.4217 | 1.3 | 0.0597271 |
| YOL140W | ARG8 | acetylornithine transaminase | 0.4225 | 1.3 | 0.0259812 |
| YKL190W | CNB1 | calcium ion binding | 0.4225 | 1.3 | 0.0636552 |
| YGR235C | | unknown | 0.4242 | 1.3 | 0.0524528 |
| YDR014W | RAD61 | unknown | 0.4247 | 1.3 | 0.0111871 |
| YDL191W | RPL35A | structural constituent of ribosome | 0.4253 | 1.3 | 0.0357698 |
| YLR462W | | unknown | 0.4260 | 1.3 | 0.0943067 |
| YBR059C | AKL1 | protein kinase | 0.4262 | 1.3 | 0.107159 |
| YGR124W | ASN2 | asparagine synthase (glutamine-hydrolyzing) | 0.4268 | 1.3 | 0.134251 |
| YLR049C | | unknown | 0.4270 | 1.3 | 0.0305927 |
| YGR283C | | unknown | 0.4275 | 1.3 | 0.0132946 |
| YOL071W | EMI5 | unknown | 0.4280 | 1.3 | 0.0477771 |
| YLR225C | | unknown | 0.4285 | 1.3 | 0.132513 |
| YCR036W | RBK1 | ATP binding | 0.4285 | 1.3 | 0.0424718 |
| YKL143W | LTV1 | unknown | 0.4312 | 1.3 | 0.0801295 |
| YMR149W | SWP1 | dolichyl-diphosphooligosaccharide-protein glycotransferase | 0.4323 | 1.3 | 0.026648 |
| YML070W | DAK1 | glycerone kinase | 0.4327 | 1.3 | 0.224696 |
| YDR344C | | | 0.4328 | 1.3 | 0.174276 |
| YML108W | | unknown | 0.4332 | 1.4 | 0.114318 |
| YPL159C | PET20 | unknown | 0.4343 | 1.4 | 0.128292 |
| YPL112C | PEX25 | unknown | 0.4355 | 1.4 | 0.119415 |
| YKL107W | | unknown | 0.4367 | 1.4 | 0.0381975 |
| YHR021C | RPS27B | structural constituent of ribosome | 0.4367 | 1.4 | 0.0537606 |
| YFL034C-B | MOB2 | protein kinase | 0.4368 | 1.4 | 0.0579427 |
| YDL145C | COP1 | unknown | 0.4373 | 1.4 | 0.106422 |
| YLR257W | | unknown | 0.4380 | 1.4 | 0.00559337 |
| YDL156W | | unknown | 0.4385 | 1.4 | 0.0119992 |
| YMR050C | | | 0.4388 | 1.4 | 0.036899 |
| YGL187C | COX4 | cytochrome-c oxidase | 0.4388 | 1.4 | 0.0877599 |
| YMR021C | MAC1 | specific RNA polymerase II transcription factor | 0.4390 | 1.4 | 0.206785 |
| YDR225W | HTA1 | DNA binding | 0.4393 | 1.4 | 0.161979 |
| YGR246C | BRF1 | RNA polymerase III transcription factor | 0.4393 | 1.4 | 0.168143 |
| YJL216C | | alpha-galactosidase * | 0.4400 | 1.4 | 0.0218329 |
| YDR515W | SLF1 | RNA binding | 0.4403 | 1.4 | 0.0740588 |
| YJL031C | BET4 | Rab-protein geranylgeranyltransferase | 0.4415 | 1.4 | 0.00258431 |
| YOR023C | AHC1 | histone acetyltransferase | 0.4425 | 1.4 | 0.205194 |
| YGR095C | RRP46 | 3'-5'-exoribonuclease | 0.4432 | 1.4 | 0.086252 |
| YEL050C | RML2 | structural constituent of ribosome | 0.4433 | 1.4 | 0.0341233 |
| YLR204W | QRI5 | unknown | 0.4437 | 1.4 | 0.0623012 |
| YKL139W | CTK1 | protein kinase | 0.4438 | 1.4 | 0.198831 |
| YMR173W | DDR48 | ATPase * | 0.4440 | 1.4 | 0.0711408 |
| YHR214W-A | | | 0.4442 | 1.4 | 0.104409 |
| YDR367W | | unknown | 0.4447 | 1.4 | 0.0351575 |
| YNR061C | | unknown | 0.4450 | 1.4 | 0.0823606 |
| YDL008W | APC11 | protein binding | 0.4460 | 1.4 | 0.00663904 |
| YFL018W-A | | | 0.4468 | 1.4 | 0.0780435 |
| YDR210W-A | | | 0.4472 | 1.4 | 0.000477681 |
| YOR254C | SEC63 | protein transporter | 0.4473 | 1.4 | 0.17585 |
| YDR027C | VPS54 | unknown | 0.4475 | 1.4 | 0.0846015 |
| YHR100C | | unknown | 0.4477 | 1.4 | 0.0133828 |
| YOL139C | CDC33 | translation initiation factor | 0.4477 | 1.4 | 0.0221916 |
| YOR222W | ODC2 | intracellular transporter | 0.4477 | 1.4 | 0.311996 |
| YML129C | COX14 | unknown | 0.4492 | 1.4 | 0.0677092 |
| YMR012W | CLU1 | unknown | 0.4492 | 1.4 | 0.254401 |
| YGR147C | NAT2 | peptide alpha-N-acetyltransferase | 0.4493 | 1.4 | 0.0441321 |
| YBR172C | SMY2 | unknown | 0.4497 | 1.4 | 0.0108304 |
| YMR177W | MMT1 | unknown | 0.4497 | 1.4 | 0.0136465 |
| YIL076W | SEC28 | unknown | 0.4498 | 1.4 | 0.00542584 |
| YLR250W | SSP120 | unknown | 0.4500 | 1.4 | 0.0343701 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-------------|
| YOR248W | | | 0.4500 | 1.4 | 0.129292 |
| YKR100C | SKG1 | unknown | 0.4515 | 1.4 | 0.272004 |
| YHR103W | SBE22 | unknown | 0.4517 | 1.4 | 0.13089 |
| YGR093W | | unknown | 0.4518 | 1.4 | 0.0218532 |
| YDR216W | ADR1 | transcription factor | 0.4523 | 1.4 | 0.0985205 |
| YGL129C | RSM23 | structural constituent of ribosome | 0.4525 | 1.4 | 0.13619 |
| YHL013C | OTU2 | unknown | 0.4528 | 1.4 | 0.0860462 |
| YPL128C | TBF1 | DNA binding | 0.4533 | 1.4 | 0.204133 |
| YJR055W | HIT1 | unknown | 0.4538 | 1.4 | 0.287507 |
| YGR202C | PCT1 | choline-phosphate cytidyltransferase | 0.4540 | 1.4 | 0.310226 |
| YPL237W | SUI3 | translation initiation factor | 0.4550 | 1.4 | 0.0280956 |
| YIL072W | HOP1 | DNA binding | 0.4562 | 1.4 | 0.073242 |
| YPR120C | CLB5 | cyclin-dependent protein kinase | 0.4570 | 1.4 | 0.0899671 |
| YGL219C | MDM34 | unknown | 0.4575 | 1.4 | 0.039917 |
| YGR080W | TWF1 | actin monomer binding | 0.4575 | 1.4 | 0.0650006 |
| YHL006C | SHU1 | unknown | 0.4578 | 1.4 | 0.101686 |
| YDL054C | MCH1 | transporter | 0.4587 | 1.4 | 0.121718 |
| YMR228W | MTF1 | S-adenosylmethionine-dependent methyltransferase * | 0.4592 | 1.4 | 0.146672 |
| YDR486C | VPS60 | unknown | 0.4595 | 1.4 | 0.00029335 |
| YMR051C | | | 0.4597 | 1.4 | 0.250944 |
| YDR085C | AFR1 | receptor signaling protein | 0.4610 | 1.4 | 0.158499 |
| YGR257C | MTM1 | transporter | 0.4613 | 1.4 | 0.0341731 |
| YHR186C | KOG1 | unknown | 0.4615 | 1.4 | 0.0298282 |
| YBL064C | PRX1 | thioredoxin peroxidase | 0.4622 | 1.4 | 0.0455349 |
| YJR032W | CPR7 | unfolded protein binding | 0.4622 | 1.4 | 0.0764775 |
| YER182W | FMP10 | unknown | 0.4633 | 1.4 | 0.0834156 |
| YJL166W | QCR8 | ubiquinol-cytochrome-c reductase | 0.4633 | 1.4 | 0.0199569 |
| YOR182C | RPS30B | structural constituent of ribosome | 0.4637 | 1.4 | 0.0422405 |
| YDL147W | RPN5 | unknown | 0.4640 | 1.4 | 0.143632 |
| YFR052W | RPN12 | endopeptidase | 0.4657 | 1.4 | 0.0550977 |
| YLR354C | TAL1 | transaldolase | 0.4657 | 1.4 | 0.0421182 |
| YKL151C | | unknown | 0.4660 | 1.4 | 0.0270596 |
| YHR121W | LSM12 | RNA binding | 0.4660 | 1.4 | 0.0620676 |
| YOR017W | PET127 | unknown | 0.4662 | 1.4 | 0.000291239 |
| YBR010W | HHT1 | DNA binding | 0.4663 | 1.4 | 0.0427297 |
| YBR021W | FUR4 | uracil permease | 0.4683 | 1.4 | 0.00678431 |
| YLR252W | | | 0.4697 | 1.4 | 0.0312779 |
| YGL057C | | unknown | 0.4700 | 1.4 | 0.160958 |
| YDR031W | | unknown | 0.4703 | 1.4 | 0.0218881 |
| YJL151C | SNA3 | unknown | 0.4708 | 1.4 | 0.010204 |
| YER134C | | unknown | 0.4712 | 1.4 | 0.0681911 |
| YLR410W-A | | | 0.4718 | 1.4 | 0.157633 |
| YJR080C | FMP26 | unknown | 0.4718 | 1.4 | 0.0411835 |
| YPR040W | TIP41 | unknown | 0.4722 | 1.4 | 0.00659972 |
| YER028C | MIG3 | DNA binding | 0.4727 | 1.4 | 0.138613 |
| YHR167W | THP2 | nucleic acid binding | 0.4727 | 1.4 | 0.135413 |
| YDL101C | DUN1 | protein kinase | 0.4728 | 1.4 | 0.0072843 |
| YIL056W | VHR1 | DNA binding | 0.4740 | 1.4 | 0.178737 |
| YNL131W | TOM22 | protein transporter | 0.4740 | 1.4 | 0.0189604 |
| YOR260W | GCD1 | translation initiation factor | 0.4748 | 1.4 | 0.0792931 |
| YJR010W | MET3 | sulfate adenylyltransferase (ATP) | 0.4767 | 1.4 | 0.00783133 |
| YGR001C | | methyltransferase | 0.4768 | 1.4 | 0.166606 |
| YDR013W | PSF1 | unknown | 0.4775 | 1.4 | 0.149795 |
| YCL031C | RRP7 | unknown | 0.4783 | 1.4 | 0.00567298 |
| YGL169W | SUA5 | unknown | 0.4785 | 1.4 | 0.173407 |
| YHL009C | YAP3 | transcription factor | 0.4788 | 1.4 | 0.198653 |
| YOR174W | MED4 | RNA polymerase II transcription mediator | 0.4802 | 1.4 | 0.0712662 |
| YNL137C | NAM9 | structural constituent of ribosome | 0.4807 | 1.4 | 0.0328999 |
| YMR039C | SUB1 | transcription coactivator | 0.4823 | 1.4 | 0.0127394 |
| YNR054C | ESF2 | transcription regulator | 0.4835 | 1.4 | 0.0436859 |
| YOR087W | YVC1 | calcium channel * | 0.4835 | 1.4 | 0.0917948 |
| YJL102W | MEF2 | translation elongation factor | 0.4837 | 1.4 | 0.00754178 |
| YPL014W | | unknown | 0.4842 | 1.4 | 0.070502 |
| YIL131C | FKH1 | transcription factor | 0.4857 | 1.4 | 0.274306 |
| YLR294C | | | 0.4858 | 1.4 | 0.023396 |
| YBR037C | SCO1 | thioredoxin peroxidase | 0.4863 | 1.4 | 0.0361619 |
| YFR025C | HIS2 | histidinol-phosphatase | 0.4873 | 1.4 | 0.247822 |
| YHR051W | COX6 | cytochrome-c oxidase | 0.4878 | 1.4 | 0.0538899 |
| YMR281W | GPI12 | N-acetylglucosaminylphosphatidylinositol deacetylase | 0.4880 | 1.4 | 0.213246 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YDR281C | PHM6 | unknown | 0.4885 | 1.4 | 0.0241516 |
| YFR017C | | unknown | 0.4893 | 1.4 | 0.216405 |
| YEL066W | HPA3 | histone acetyltransferase | 0.4895 | 1.4 | 0.00210665 |
| YOR158W | PET123 | structural constituent of ribosome | 0.4897 | 1.4 | 0.104932 |
| YDR206W | EBS1 | unknown | 0.4898 | 1.4 | 0.100733 |
| YOR045W | TOM6 | protein transporter | 0.4900 | 1.4 | 0.110627 |
| YLR295C | ATP14 | hydrogen-transporting ATP synthase , rotational mechanism | 0.4915 | 1.4 | 0.0115678 |
| YBR119W | MUD1 | RNA binding | 0.4925 | 1.4 | 0.0598601 |
| YBR202W | CDC47 | chromatin binding | 0.4927 | 1.4 | 0.0580251 |
| YDR299W | BFR2 | unknown | 0.4933 | 1.4 | 0.0469495 |
| YMR073C | | unknown | 0.4942 | 1.4 | 0.0305443 |
| YBL066C | SEF1 | unknown | 0.4962 | 1.4 | 0.0157141 |
| YNR073C | | mannitol dehydrogenase | 0.4967 | 1.4 | 0.168241 |
| YPL010W | RET3 | protein binding | 0.4972 | 1.4 | 0.0574156 |
| YDR081C | PDC2 | transcription regulator | 0.4975 | 1.4 | 0.0223707 |
| YCL007C | | | 0.4975 | 1.4 | 0.174987 |
| YDL114W | | unknown | 0.4982 | 1.4 | 0.164575 |
| YLL014W | | unknown | 0.4993 | 1.4 | 0.063809 |
| YDR049W | | unknown | 0.5002 | 1.4 | 0.395127 |
| YFR003C | YPI1 | protein phosphatase inhibitor | 0.5007 | 1.4 | 0.0451772 |
| YHR083W | SAM35 | protein binding | 0.5007 | 1.4 | 0.067767 |
| YIL035C | CKA1 | protein kinase | 0.5010 | 1.4 | 0.0533489 |
| YJL063C | MRPL8 | structural constituent of ribosome | 0.5013 | 1.4 | 0.0767068 |
| YCR020C-A | MAK31 | peptide alpha-N-acetyltransferase | 0.5015 | 1.4 | 0.0163273 |
| YER127W | LCP5 | RNA binding | 0.5017 | 1.4 | 0.00670135 |
| YKL218C | SRY1 | threo-3-hydroxyaspartate ammonia-lyase * | 0.5020 | 1.4 | 0.0317813 |
| YCL006C | | | 0.5032 | 1.4 | 0.263108 |
| YDR202C | RAV2 | unknown | 0.5048 | 1.4 | 0.0162182 |
| YML127W | RSC9 | chromatin binding | 0.5050 | 1.4 | 0.183193 |
| YDR532C | | unknown | 0.5053 | 1.4 | 0.0552863 |
| YIR011C | STS1 | unknown | 0.5055 | 1.4 | 0.0311946 |
| YOR372C | NDD1 | transcriptional activator | 0.5057 | 1.4 | 0.0649134 |
| YDR529C | QCR7 | ubiquinol-cytochrome-c reductase | 0.5062 | 1.4 | 0.0913272 |
| YKL161C | | protein kinase | 0.5063 | 1.4 | 0.0101032 |
| YPL145C | KES1 | oxysterol binding | 0.5067 | 1.4 | 0.0949829 |
| YKR007W | MEH1 | unknown | 0.5067 | 1.4 | 0.0412531 |
| YGR015C | | unknown | 0.5073 | 1.4 | 0.0985756 |
| YDR318W | MCM21 | protein binding | 0.5082 | 1.4 | 0.0181467 |
| YGR013W | SNU71 | RNA binding | 0.5083 | 1.4 | 0.19209 |
| YPL091W | GLR1 | glutathione-disulfide reductase | 0.5085 | 1.4 | 0.0188051 |
| YDR332W | IRC3 | helicase | 0.5087 | 1.4 | 0.0594006 |
| YIR021W | MRS1 | RNA binding | 0.5100 | 1.4 | 0.0860763 |
| YFL043C | | | 0.5102 | 1.4 | 0.0215182 |
| YMR269W | TMA23 | unknown | 0.5103 | 1.4 | 0.149235 |
| YDR140W | MTQ2 | S-adenosylmethionine-dependent methyltransferase * | 0.5143 | 1.4 | 0.00244867 |
| YOR167C | RPS28A | structural constituent of ribosome | 0.5147 | 1.4 | 0.0109307 |
| YDR398W | UTP5 | snoRNA binding | 0.5157 | 1.4 | 0.273772 |
| YDR096W | GIS1 | transcription factor | 0.5162 | 1.4 | 0.0165738 |
| YBR039W | ATP3 | hydrogen-transporting ATP synthase , rotational mechanism | 0.5168 | 1.4 | 0.00768298 |
| YDL235C | YPD1 | transferase , transferring phosphorus-containing groups | 0.5168 | 1.4 | 0.030936 |
| YJL046W | | unknown | 0.5172 | 1.4 | 0.0786259 |
| YOR187W | TUF1 | GTPase * | 0.5173 | 1.4 | 0.0045012 |
| YBL113C | | helicase | 0.5183 | 1.4 | 0.0112937 |
| YMR174C | PAI3 | endopeptidase inhibitor | 0.5185 | 1.4 | 0.0563961 |
| YNL192W | CHS1 | chitin synthase | 0.5187 | 1.4 | 0.0346981 |
| YBR120C | CBP6 | unknown | 0.5190 | 1.4 | 0.105301 |
| YGL093W | SPC105 | structural constituent of cytoskeleton | 0.5193 | 1.4 | 0.040631 |
| YAL047C | SPC72 | structural constituent of cytoskeleton | 0.5195 | 1.4 | 0.0531787 |
| YDL048C | STP4 | unknown | 0.5198 | 1.4 | 0.000386249 |
| YJL007C | | | 0.5200 | 1.4 | 0.0460419 |
| YGL242C | | unknown | 0.5203 | 1.4 | 0.0888156 |
| YDR117C | TMA64 | RNA binding | 0.5215 | 1.4 | 0.348711 |
| YCL043C | PDI1 | protein disulfide isomerase | 0.5220 | 1.4 | 0.00948149 |
| YIL162W | SUC2 | beta-fructofuranosidase | 0.5223 | 1.4 | 0.00674801 |
| YLR179C | | unknown | 0.5223 | 1.4 | 0.0161588 |
| YNR012W | URK1 | uridine kinase | 0.5228 | 1.4 | 0.0799909 |
| YPL092W | SSU1 | sulfite transporter | 0.5230 | 1.4 | 0.0372879 |
| YNL310C | ZIM17 | protein binding | 0.5232 | 1.4 | 0.0147868 |
| YKL183W | LOT5 | unknown | 0.5233 | 1.4 | 0.146996 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-------------|
| YDL120W | YFH1 | enzyme regulator | 0.5235 | 1.4 | 0.0262338 |
| YOR046C | DBP5 | RNA helicase | 0.5240 | 1.4 | 0.194869 |
| YBR015C | MNN2 | alpha-1,2-mannosyltransferase | 0.5242 | 1.4 | 0.0328329 |
| YLR050C | | unknown | 0.5253 | 1.4 | 0.0285192 |
| YCR039C | MATALPHA2 | transcription corepressor | 0.5260 | 1.4 | 0.0609649 |
| YOR062C | | unknown | 0.5262 | 1.4 | 0.0580951 |
| YPR041W | TIF5 | translation initiation factor | 0.5263 | 1.4 | 0.0235665 |
| YKL133C | | unknown | 0.5268 | 1.4 | 0.146588 |
| YDR252W | BTT1 | unfolded protein binding | 0.5297 | 1.4 | 0.221855 |
| YKR030W | GMH1 | unknown | 0.5297 | 1.4 | 0.155926 |
| YOR236W | DFR1 | dihydrofolate reductase | 0.5302 | 1.4 | 0.0470777 |
| YKL170W | MRPL38 | structural constituent of ribosome | 0.5310 | 1.4 | 0.0147879 |
| YDR002W | YRB1 | Ran GTPase binding | 0.5318 | 1.4 | 0.00305125 |
| YAL060W | BDH1 | (R,R)-butanediol dehydrogenase | 0.5327 | 1.4 | 0.0479707 |
| YBR141C | | unknown | 0.5335 | 1.4 | 0.00282584 |
| YNL281W | HCH1 | chaperone activator | 0.5338 | 1.4 | 0.057266 |
| YOR297C | TIM18 | protein transporter | 0.5348 | 1.4 | 0.0431008 |
| YOR208W | PTP2 | protein tyrosine phosphatase | 0.5352 | 1.4 | 0.0784979 |
| YMR292W | GOT1 | unknown | 0.5355 | 1.4 | 0.0436418 |
| TMR307W | | | 0.5367 | 1.5 | 0.027773 |
| YLR023C | IZH3 | metal ion binding | 0.5367 | 1.5 | 0.0859464 |
| YLR053C | | unknown | 0.5372 | 1.5 | 0.042214 |
| YDR530C | APA2 | bis(5'-nucleosyl)-tetraphosphatase * | 0.5373 | 1.5 | 0.0556706 |
| YHR040W | BCD1 | nucleic acid binding | 0.5383 | 1.5 | 0.00665666 |
| YBL002W | HTB2 | DNA binding | 0.5383 | 1.5 | 0.0273166 |
| YDR412W | RRP17 | unknown | 0.5385 | 1.5 | 0.0622606 |
| YOL137W | BSC6 | unknown | 0.5387 | 1.5 | 0.0190231 |
| YNL132W | KRE33 | unknown | 0.5392 | 1.5 | 0.0706516 |
| YGL263W | COS12 | unknown | 0.5393 | 1.5 | 0.046386 |
| YDL157C | | unknown | 0.5410 | 1.5 | 0.0447279 |
| YBL090W | MRP21 | structural constituent of ribosome | 0.5412 | 1.5 | 0.0317772 |
| YJR077C | MIR1 | inorganic phosphate transporter | 0.5415 | 1.5 | 0.00792206 |
| YHR137W | ARO9 | aromatic-amino-acid transaminase | 0.5423 | 1.5 | 0.0176031 |
| YBR091C | MRS5 | protein transporter | 0.5425 | 1.5 | 0.0726096 |
| YLR300W | EXG1 | glucan 1,3-beta-glucosidase | 0.5438 | 1.5 | 0.160733 |
| YBR070C | ALG14 | N-acetylglucosaminyldiphosphodichol N-acetylglucosam | 0.5440 | 1.5 | 0.00624406 |
| YGL163C | RAD54 | DNA-dependent ATPase * | 0.5443 | 1.5 | 0.0598154 |
| YCR080W | | | 0.5445 | 1.5 | 0.136618 |
| YDR331W | GPI8 | GPI-anchor transamidase | 0.5447 | 1.5 | 0.105054 |
| YKL003C | MRP17 | structural constituent of ribosome | 0.5450 | 1.5 | 0.0536721 |
| YLR016C | PML1 | unknown | 0.5475 | 1.5 | 0.0068785 |
| YEL026W | SNU13 | RNA binding | 0.5477 | 1.5 | 0.0559199 |
| YNL217W | | phosphoric monoester hydrolase | 0.5477 | 1.5 | 0.000190487 |
| YPR100W | MRPL51 | structural constituent of ribosome | 0.5478 | 1.5 | 0.00524761 |
| YNR017W | MAS6 | protein transporter | 0.5478 | 1.5 | 0.0219246 |
| YDL241W | | unknown | 0.5480 | 1.5 | 0.0436493 |
| YMR048W | CSM3 | unknown | 0.5490 | 1.5 | 0.00704276 |
| YDL069C | CBS1 | translation factor | 0.5505 | 1.5 | 0.0490582 |
| YFR011C | | unknown | 0.5508 | 1.5 | 0.0310636 |
| YJR136C | | unknown | 0.5530 | 1.5 | 0.118162 |
| YGR062C | COX18 | protein transporter | 0.5533 | 1.5 | 0.113301 |
| YBR030W | | unknown | 0.5538 | 1.5 | 0.108062 |
| YDL092W | SRP14 | signal sequence binding | 0.5543 | 1.5 | 0.0537937 |
| YIL040W | APQ12 | unknown | 0.5550 | 1.5 | 0.129801 |
| YGL107C | RMD9 | unknown | 0.5550 | 1.5 | 0.122977 |
| YFL-TYA | | | 0.5557 | 1.5 | 0.094898 |
| YEL003W | GIM4 | tubulin binding | 0.5563 | 1.5 | 0.00555931 |
| YNL031C | HHT2 | DNA binding | 0.5580 | 1.5 | 0.0133833 |
| YDR462W | MRPL28 | structural constituent of ribosome | 0.5590 | 1.5 | 0.00962826 |
| YDR273W | DON1 | unknown | 0.5603 | 1.5 | 0.00170895 |
| YLR192C | HCR1 | translation initiation factor | 0.5603 | 1.5 | 0.00171666 |
| YER150W | SP11 | unknown | 0.5608 | 1.5 | 0.00880508 |
| YCL026C | | | 0.5612 | 1.5 | 0.0377089 |
| YCL044C | MGR1 | unknown | 0.5612 | 1.5 | 0.00251144 |
| YNL036W | NCE103 | carbonate dehydratase | 0.5613 | 1.5 | 0.000695624 |
| YDR540C | IRC4 | unknown | 0.5622 | 1.5 | 0.170174 |
| YFL010C | WWM1 | unknown | 0.5642 | 1.5 | 0.00405003 |
| YOR305W | | unknown | 0.5653 | 1.5 | 0.0456689 |
| YNR036C | | structural constituent of ribosome | 0.5657 | 1.5 | 0.0325287 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|------------|
| YBR171W | SEC66 | protein transporter | 0.5658 | 1.5 | 0.0532421 |
| YKR011C | | unknown | 0.5683 | 1.5 | 0.165897 |
| YDR187C | | | 0.5683 | 1.5 | 0.00615473 |
| YDL089W | | unknown | 0.5683 | 1.5 | 0.0411234 |
| YFL035C-A | | | 0.5687 | 1.5 | 0.245016 |
| YBR170C | NPL4 | unknown | 0.5697 | 1.5 | 0.0377509 |
| YKL077W | | unknown | 0.5743 | 1.5 | 0.00426023 |
| YIL063C | YRB2 | unknown | 0.5758 | 1.5 | 0.00334562 |
| YER070W | RNR1 | ribonucleoside-diphosphate reductase | 0.5758 | 1.5 | 0.0390104 |
| YDR378C | LSM6 | RNA binding | 0.5778 | 1.5 | 0.00871113 |
| YJL097W | PHS1 | unknown | 0.5778 | 1.5 | 0.00374897 |
| YFL041W | FET5 | ferroxidase | 0.5793 | 1.5 | 0.0528921 |
| YJL221C | FSP2 | alpha-glucosidase | 0.5795 | 1.5 | 0.00854282 |
| YPR094W | RDS3 | unknown | 0.5797 | 1.5 | 0.148855 |
| YIL122W | POG1 | specific RNA polymerase II transcription factor | 0.5798 | 1.5 | 0.0537962 |
| YLR251W | SYM1 | unknown | 0.5802 | 1.5 | 0.070233 |
| YDR268W | MSW1 | tryptophan-tRNA ligase | 0.5805 | 1.5 | 0.0589518 |
| YOL059W | GPD2 | glycerol-3-phosphate dehydrogenase | 0.5818 | 1.5 | 0.0497565 |
| YPL204W | HRR25 | casein kinase | 0.5840 | 1.5 | 0.0151331 |
| YJL062W | LAS21 | transferase | 0.5850 | 1.5 | 0.0290046 |
| YIL008W | URM1 | protein tag | 0.5850 | 1.5 | 0.128733 |
| YCR096C | HMRA2 | unknown | 0.5860 | 1.5 | 0.00763131 |
| YHR081W | LRP1 | unknown | 0.5863 | 1.5 | 0.0419276 |
| YAL069W | | | 0.5863 | 1.5 | 0.0812503 |
| YLR230W | | | 0.5890 | 1.5 | 0.606237 |
| YOL011W | PLB3 | lysophospholipase | 0.5898 | 1.5 | 0.0195474 |
| YOR016C | ERP4 | unknown | 0.5913 | 1.5 | 0.0772978 |
| YDR235W | PRP42 | RNA binding | 0.5913 | 1.5 | 0.0061686 |
| YDL111C | RRP42 | 3'-5'-exoribonuclease | 0.5913 | 1.5 | 0.0613521 |
| YJR058C | APS2 | unknown | 0.5913 | 1.5 | 0.0135534 |
| YBL107C | | unknown | 0.5920 | 1.5 | 0.0052672 |
| Q0120 | BI4 | RNA binding | 0.5930 | 1.5 | 0.108355 |
| YGR078C | PAC10 | tubulin binding | 0.5932 | 1.5 | 0.0493755 |
| YOR338W | | unknown | 0.5937 | 1.5 | 0.130278 |
| YJL208C | NUC1 | ribonuclease * | 0.5945 | 1.5 | 0.0805223 |
| Q0255 | | unknown | 0.5947 | 1.5 | 0.0107965 |
| YOR201C | MRM1 | rRNA (guanine-N1-)-methyltransferase | 0.5947 | 1.5 | 0.0781559 |
| YDR517W | GRH1 | unknown | 0.5960 | 1.5 | 0.00537524 |
| YMR173W-A | | | 0.5963 | 1.5 | 0.0421073 |
| YDR092W | UBC13 | ubiquitin conjugating enzyme | 0.5977 | 1.5 | 0.012112 |
| YJL148W | RPA34 | DNA-directed RNA polymerase | 0.5980 | 1.5 | 0.0328234 |
| YDL017W | CDC7 | protein serine/threonine kinase | 0.5980 | 1.5 | 0.10662 |
| YFL016C | MDJ1 | unfolded protein binding | 0.5990 | 1.5 | 0.013336 |
| YGR172C | YIP1 | unknown | 0.5997 | 1.5 | 0.0237079 |
| YOL022C | | unknown | 0.6008 | 1.5 | 0.0614175 |
| YDL055C | PSA1 | mannose-1-phosphate guanylyltransferase | 0.6012 | 1.5 | 0.013402 |
| YKL082C | RRP14 | unknown | 0.6023 | 1.5 | 0.0951916 |
| YPL106C | SSE1 | ATP binding | 0.6028 | 1.5 | 0.00288947 |
| YKL150W | MCR1 | cytochrome-b5 reductase | 0.6030 | 1.5 | 0.00263223 |
| YNL177C | MRPL22 | structural constituent of ribosome | 0.6030 | 1.5 | 0.0188789 |
| YDR290W | | | 0.6043 | 1.5 | 0.189285 |
| YMR290C | HAS1 | RNA binding | 0.6045 | 1.5 | 0.0269443 |
| YBR182C | SMP1 | transcription factor | 0.6052 | 1.5 | 0.0141909 |
| YHR147C | MRPL6 | structural constituent of ribosome | 0.6060 | 1.5 | 0.0299516 |
| YGL202W | ARO8 | aromatic-amino-acid transaminase | 0.6070 | 1.5 | 0.167251 |
| YLL011W | SOF1 | snoRNA binding | 0.6078 | 1.5 | 0.0308592 |
| YPL148C | PPT2 | phosphopantetheinyltransferase | 0.6082 | 1.5 | 0.0654919 |
| YOR306C | MCH5 | transporter | 0.6083 | 1.5 | 0.104411 |
| YCLX01W | | | 0.6093 | 1.5 | 0.0203037 |
| YLR186W | EMG1 | unknown | 0.6097 | 1.5 | 0.0908688 |
| YML125C | PGA3 | unknown | 0.6128 | 1.5 | 0.0396362 |
| YNL306W | MRPS18 | structural constituent of ribosome | 0.6142 | 1.5 | 0.0253688 |
| YHR180W | | | 0.6148 | 1.5 | 0.068993 |
| YJL137C | GLG2 | glycogenin glucosyltransferase | 0.6148 | 1.5 | 0.025975 |
| YDR028C | REG1 | protein phosphatase type 1 regulator | 0.6152 | 1.5 | 0.00565826 |
| YKL011C | CCE1 | endodeoxyribonuclease | 0.6155 | 1.5 | 0.105211 |
| YGR002C | SWC4 | DNA binding | 0.6160 | 1.5 | 0.103682 |
| YEL001C | | unknown | 0.6160 | 1.5 | 0.0063596 |
| YOL005C | RPB11 | DNA-directed RNA polymerase | 0.6168 | 1.5 | 0.0163654 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|------------|
| YBR173C | UMP1 | proteasome activator | 0.6175 | 1.5 | 0.0240801 |
| YLR275W | SMD2 | RNA splicing factor | 0.6180 | 1.5 | 0.0961118 |
| YCR027C | RHB1 | GTPase | 0.6185 | 1.5 | 0.0921225 |
| YKL089W | MIF2 | centromeric DNA binding | 0.6188 | 1.5 | 0.0108698 |
| YGR156W | PTI1 | RNA binding | 0.6193 | 1.5 | 0.143618 |
| YKL035W | UGP1 | UTP:glucose-1-phosphate uridylyltransferase | 0.6195 | 1.5 | 0.0231547 |
| YDR044W | HEM13 | coproporphyrinogen oxidase | 0.6198 | 1.5 | 0.0148295 |
| YDR079W | PET100 | unfolded protein binding | 0.6203 | 1.5 | 0.0273947 |
| YGL070C | RPB9 | DNA-directed RNA polymerase | 0.6208 | 1.5 | 0.0836544 |
| Q0250 | COX2 | cytochrome-c oxidase | 0.6220 | 1.5 | 0.033301 |
| YCL067C | HMLALPHA2 | transcription corepressor | 0.6222 | 1.5 | 0.0443276 |
| YGR076C | MRPL25 | structural constituent of ribosome | 0.6225 | 1.5 | 0.070643 |
| YGR020C | VMA7 | hydrogen-transporting ATPase , rotational mechanism | 0.6230 | 1.5 | 0.00698579 |
| YOL023W | IFM1 | RNA binding | 0.6230 | 1.5 | 0.0737444 |
| YML078W | CPR3 | peptidyl-prolyl cis-trans isomerase | 0.6233 | 1.5 | 0.0494883 |
| YDR489W | SLD5 | unknown | 0.6238 | 1.5 | 0.00428904 |
| YMR093W | UTP15 | snoRNA binding | 0.6242 | 1.5 | 0.109977 |
| YAL007C | ERP2 | unknown | 0.6262 | 1.5 | 0.0525176 |
| YDR373W | FRQ1 | enzyme activator * | 0.6278 | 1.5 | 0.00456051 |
| YFL032W | | | 0.6278 | 1.5 | 0.127398 |
| YGL226W | | unknown | 0.6282 | 1.5 | 0.00366272 |
| YGR152C | RSR1 | GTPase * | 0.6283 | 1.5 | 0.0115137 |
| YGL143C | MRF1 | translation release factor | 0.6288 | 1.5 | 0.0788082 |
| YNL153C | GIM3 | tubulin binding | 0.6293 | 1.5 | 0.109134 |
| YKL225W | | | 0.6298 | 1.5 | 0.0665314 |
| YNL100W | | unknown | 0.6303 | 1.5 | 0.0180901 |
| YPL271W | ATP15 | hydrogen-transporting ATP synthase , rotational mechanism | 0.6348 | 1.6 | 0.012649 |
| YHR122W | | unknown | 0.6380 | 1.6 | 0.0317915 |
| YDR173C | ARG82 | inositol or phosphatidylinositol kinase | 0.6392 | 1.6 | 0.0163209 |
| YLR202C | | | 0.6393 | 1.6 | 0.0496014 |
| YCR034W | FEN1 | fatty acid elongase | 0.6393 | 1.6 | 0.0178405 |
| YPL215W | CBP3 | unknown | 0.6397 | 1.6 | 0.0830257 |
| YOR190W | SPR1 | glucan 1,3-beta-glucosidase | 0.6398 | 1.6 | 0.0863526 |
| YJL196C | ELO1 | fatty acid elongase | 0.6432 | 1.6 | 0.100051 |
| YBR267W | REI1 | unknown | 0.6462 | 1.6 | 0.0318328 |
| YPR176C | BET2 | Rab-protein geranylgeranyltransferase | 0.6500 | 1.6 | 0.293145 |
| YDR066C | | unknown | 0.6500 | 1.6 | 0.202496 |
| YJL104W | PAM16 | protein binding | 0.6505 | 1.6 | 0.03308 |
| YJL054W | TIM54 | protein transporter | 0.6515 | 1.6 | 0.0767007 |
| YCL035C | GRX1 | thiol-disulfide exchange intermediate * | 0.6542 | 1.6 | 0.00238789 |
| YHR116W | COX23 | unknown | 0.6558 | 1.6 | 0.0277786 |
| YDL121C | | unknown | 0.6572 | 1.6 | 0.0951637 |
| YGR063C | SPT4 | RNA polymerase II transcription elongation factor | 0.6647 | 1.6 | 0.13183 |
| YOR354C | MSC6 | unknown | 0.6647 | 1.6 | 0.145648 |
| YKL016C | ATP7 | structural molecule * | 0.6663 | 1.6 | 0.0258672 |
| YKL085W | MDH1 | L-malate dehydrogenase | 0.6663 | 1.6 | 0.00284255 |
| YNL108C | | unknown | 0.6665 | 1.6 | 0.107433 |
| YJR156C | THI11 | protein binding | 0.6675 | 1.6 | 0.0200628 |
| YLR221C | RSA3 | unknown | 0.6687 | 1.6 | 0.129387 |
| YDR379W | RGA2 | signal transducer * | 0.6688 | 1.6 | 0.0701096 |
| YOL136C | PFK27 | 6-phosphofructo-2-kinase | 0.6692 | 1.6 | 0.0943872 |
| YDL107W | MSS2 | protein translocase | 0.6700 | 1.6 | 0.0857198 |
| YFL060C | SNO3 | unknown | 0.6703 | 1.6 | 0.00477158 |
| YBL038W | MRPL16 | structural constituent of ribosome* | 0.6707 | 1.6 | 0.0164289 |
| YPR166C | MRP2 | structural constituent of ribosome | 0.6720 | 1.6 | 0.0525754 |
| YOR279C | RFM1 | unfolded protein binding | 0.6730 | 1.6 | 0.0121076 |
| YGL043W | DST1 | positive transcription elongation factor | 0.6740 | 1.6 | 0.0338224 |
| YDR377W | ATP17 | hydrogen-transporting ATP synthase , rotational mechanism | 0.6753 | 1.6 | 0.0351545 |
| YBR251W | MRPS5 | structural constituent of ribosome | 0.6758 | 1.6 | 0.0865782 |
| YGR159C | NSR1 | RNA binding | 0.6760 | 1.6 | 0.0291377 |
| YKL023W | | unknown | 0.6767 | 1.6 | 0.0657252 |
| YKR023W | | unknown | 0.6777 | 1.6 | 0.00906859 |
| YJL114W | | | 0.6787 | 1.6 | 0.00464335 |
| YOR198C | BFR1 | RNA binding | 0.6788 | 1.6 | 0.00379246 |
| YNL312W | RFA2 | DNA binding | 0.6792 | 1.6 | 0.0128626 |
| YKR085C | MRPL20 | structural constituent of ribosome | 0.6812 | 1.6 | 0.0217817 |
| YOR247W | SRL1 | unknown | 0.6815 | 1.6 | 0.0164093 |
| YGL098W | USE1 | SNAP receptor | 0.6815 | 1.6 | 0.0690786 |
| YFR007W | | unknown | 0.6832 | 1.6 | 0.159109 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|------------|---|----------------------------|-------------|-------------|
| YER050C | RSM18 | structural constituent of ribosome | 0.6832 | 1.6 | 0.0146155 |
| YNR037C | RSM19 | structural constituent of ribosome | 0.6837 | 1.6 | 0.0330487 |
| YPL001W | HAT1 | H3/H4 histone acetyltransferase | 0.6838 | 1.6 | 0.250496 |
| YGL136C | MRM2 | rRNA (uridine-2'-O-)-methyltransferase | 0.6842 | 1.6 | 0.0102949 |
| YDR472W | TRS31 | unknown | 0.6843 | 1.6 | 0.0526128 |
| YOR074C | CDC21 | thymidylate synthase | 0.6845 | 1.6 | 0.129474 |
| YHR176W | FMO1 | monooxygenase | 0.6858 | 1.6 | 0.109132 |
| YOR210W | RPB10 | DNA-directed RNA polymerase | 0.6888 | 1.6 | 0.00796331 |
| YML039W | | | 0.6890 | 1.6 | 0.19758 |
| YIL098C | FMC1 | unknown | 0.6898 | 1.6 | 0.0174371 |
| YGR260W | TNA1 | nicotinamide mononucleotide permease | 0.6918 | 1.6 | 0.0583647 |
| YPR113W | PIS1 | CDP-diacylglycerol-inositol 3-phosphatidyltransferase | 0.6928 | 1.6 | 0.0794115 |
| YKL109W | HAP4 | transcriptional activator | 0.6950 | 1.6 | 0.0191707 |
| YOL002C | IZH2 | metal ion binding | 0.6982 | 1.6 | 0.214678 |
| YML130C | ERO1 | electron carrier | 0.6985 | 1.6 | 0.030604 |
| YBR268W | MRPL37 | structural constituent of ribosome | 0.7025 | 1.6 | 0.006982 |
| YLR271W | | unknown | 0.7032 | 1.6 | 0.0312596 |
| YGL232W | TAN1 | RNA binding | 0.7038 | 1.6 | 0.0598313 |
| YJR059W | PTK2 | protein kinase | 0.7043 | 1.6 | 0.0177033 |
| YDR320C | SWA2 | protein binding | 0.7060 | 1.6 | 0.0159393 |
| YDR476C | | unknown | 0.7062 | 1.6 | 0.0129401 |
| YBL043W | ECM13 | unknown | 0.7070 | 1.6 | 0.00649798 |
| YLR217W | | | 0.7072 | 1.6 | 0.0232707 |
| YFR053C | HXK1 | hexokinase | 0.7078 | 1.6 | 0.140574 |
| YKL024C | URA6 | uridine kinase | 0.7100 | 1.6 | 0.0826326 |
| YPR073C | LTP1 | protein tyrosine phosphatase | 0.7115 | 1.6 | 0.0554634 |
| YDR200C | VPS64 | unknown | 0.7142 | 1.6 | 0.00860777 |
| YJL184W | GON7 | unknown | 0.7147 | 1.6 | 0.000183365 |
| YOR271C | | unknown | 0.7163 | 1.6 | 0.00807593 |
| YEL052W | AFG1 | ATPase | 0.7172 | 1.6 | 0.0761875 |
| YBL018C | POP8 | ribonuclease P * | 0.7192 | 1.6 | 0.0364137 |
| YJR063W | RPA12 | DNA-directed RNA polymerase | 0.7195 | 1.6 | 0.173241 |
| YHR112C | | cystathionine beta-lyase | 0.7202 | 1.6 | 0.0867615 |
| YFL012W | | unknown | 0.7215 | 1.6 | 0.0545161 |
| YER177W | BMH1 | protein binding | 0.7215 | 1.6 | 0.00880142 |
| YNL005C | MRP7 | structural constituent of ribosome* | 0.7218 | 1.6 | 0.00438703 |
| YGL200C | EMP24 | unknown | 0.7252 | 1.7 | 0.0106196 |
| YER083C | GET2 | unknown | 0.7253 | 1.7 | 0.0231585 |
| YDR380W | ARO10 | pyruvate decarboxylase | 0.7260 | 1.7 | 0.0498865 |
| YOR298W | MUM3 | acyltransferase | 0.7263 | 1.7 | 0.0366055 |
| YDR510W | SMT3 | protein tag | 0.7270 | 1.7 | 0.0154042 |
| YNL107W | YAF9 | unknown | 0.7288 | 1.7 | 0.181542 |
| YJL174W | KRE9 | unknown | 0.7300 | 1.7 | 0.00546207 |
| YJR117W | STE24 | metalloendopeptidase * | 0.7310 | 1.7 | 0.0617768 |
| YDL100C | GET3 | ATPase | 0.7325 | 1.7 | 0.0100448 |
| YDR241W | | | 0.7332 | 1.7 | 0.00950221 |
| YBR162W-A | YSY6 | unknown | 0.7375 | 1.7 | 0.0282321 |
| YPR074C | TKL1 | transketolase | 0.7388 | 1.7 | 0.252939 |
| YPR020W | ATP20 | structural molecule | 0.7392 | 1.7 | 0.0175005 |
| YOR344C | TYE7 | transcription factor | 0.7395 | 1.7 | 0.0535386 |
| YGR048W | UFD1 | protein binding | 0.7405 | 1.7 | 0.275455 |
| YKL192C | ACP1 | acyl carrier | 0.7415 | 1.7 | 0.00593254 |
| YDR494W | RSM28 | structural constituent of ribosome | 0.7422 | 1.7 | 0.0147008 |
| YLR239C | LIP2 | ligase | 0.7438 | 1.7 | 0.0475392 |
| YML009C | MRPL39 | structural constituent of ribosome | 0.7442 | 1.7 | 0.0140823 |
| YFR050C | PRE4 | endopeptidase | 0.7443 | 1.7 | 0.011957 |
| YNR040W | | unknown | 0.7477 | 1.7 | 0.225664 |
| YGL089C | MF(ALPHA)2 | mating pheromone | 0.7498 | 1.7 | 0.163745 |
| YHR041C | SRB2 | RNA polymerase II transcription mediator | 0.7515 | 1.7 | 0.0963786 |
| YMR123W | PKR1 | unknown | 0.7540 | 1.7 | 0.0243693 |
| YIL074C | SER33 | phosphoglycerate dehydrogenase | 0.7550 | 1.7 | 0.00306486 |
| YDR260C | SWM1 | unknown | 0.7555 | 1.7 | 0.00232931 |
| YLR051C | FCF2 | unknown | 0.7563 | 1.7 | 0.0350492 |
| YDL022W | GPD1 | glycerol-3-phosphate dehydrogenase | 0.7568 | 1.7 | 0.00350797 |
| YBR114W | RAD16 | DNA-dependent ATPase | 0.7573 | 1.7 | 0.0196294 |
| YGR189C | CRH1 | unknown | 0.7573 | 1.7 | 0.00791711 |
| YOR020C | HSP10 | unfolded protein binding | 0.7615 | 1.7 | 0.000738281 |
| YJL192C | SOP4 | unknown | 0.7620 | 1.7 | 0.0103996 |
| YOR357C | SNX3 | protein binding | 0.7623 | 1.7 | 0.0238469 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YBR009C | HHF1 | DNA binding | 0.7630 | 1.7 | 0.00936158 |
| YGL088W | | | 0.7633 | 1.7 | 0.0501882 |
| YLR183C | TOS4 | transcription factor | 0.7663 | 1.7 | 0.0176643 |
| YBR262C | FMP51 | unknown | 0.7663 | 1.7 | 0.000555644 |
| YBR146W | MRPS9 | structural constituent of ribosome | 0.7673 | 1.7 | 0.00505915 |
| YMR260C | TIF11 | translation initiation factor | 0.7708 | 1.7 | 0.0117001 |
| YMR157C | FMP39 | unknown | 0.7717 | 1.7 | 0.0514704 |
| YGR183C | QCR9 | ubiquinol-cytochrome-c reductase | 0.7720 | 1.7 | 0.0335958 |
| YBR201W | DER1 | unknown | 0.7728 | 1.7 | 0.0205069 |
| YDL004W | ATP16 | hydrogen-transporting ATP synthase , rotational mechanism | 0.7730 | 1.7 | 0.00257301 |
| YGR165W | MRPS35 | structural constituent of ribosome | 0.7733 | 1.7 | 0.0154686 |
| YMR101C | SRT1 | dehydrodolichyl diphosphate synthase | 0.7735 | 1.7 | 0.000649768 |
| YDR270W | CCC2 | cation-transporting ATPase | 0.7748 | 1.7 | 0.0443133 |
| YMR188C | MRPS17 | structural constituent of ribosome | 0.7765 | 1.7 | 0.029002 |
| YJR096W | | aldo-keto reductase * | 0.7793 | 1.7 | 0.00621474 |
| YOR150W | MRPL23 | structural constituent of ribosome | 0.7797 | 1.7 | 0.0466981 |
| YGR136W | LSB1 | unknown | 0.7875 | 1.7 | 0.0084724 |
| YLR292C | SEC72 | protein transporter | 0.7902 | 1.7 | 0.0186406 |
| YDR041W | RSM10 | structural constituent of ribosome | 0.7903 | 1.7 | 0.00054655 |
| YHR128W | FUR1 | uracil phosphoribosyltransferase | 0.7942 | 1.7 | 0.0212612 |
| YDR353W | TRR1 | thioredoxin-disulfide reductase | 0.7947 | 1.7 | 0.0193076 |
| YOR224C | RPB8 | DNA-directed RNA polymerase | 0.7970 | 1.7 | 0.0757633 |
| YML004C | GLO1 | lactoylgutathione lyase | 0.7982 | 1.7 | 5.85055E-05 |
| YLR391W | | | 0.7988 | 1.7 | 0.024883 |
| YBR085W | AAC3 | ATP:ADP antiporter | 0.7993 | 1.7 | 0.0241274 |
| YDR493W | FMP36 | unknown | 0.8005 | 1.7 | 0.0537485 |
| YDR113C | PDS1 | protein binding | 0.8012 | 1.7 | 0.00346949 |
| YDR220C | | | 0.8030 | 1.7 | 0.0218942 |
| YCL051W | LRE1 | transcription regulator | 0.8053 | 1.7 | 0.0044093 |
| YLR203C | MSS51 | unknown | 0.8083 | 1.8 | 0.00957611 |
| YBL021C | HAP3 | transcriptional activator | 0.8097 | 1.8 | 0.0113595 |
| YDR151C | CTH1 | transcription factor | 0.8100 | 1.8 | 0.00504152 |
| YDL021W | GPM2 | unknown | 0.8102 | 1.8 | 0.00337149 |
| YHR034C | PIH1 | unknown | 0.8110 | 1.8 | 0.0174512 |
| YCR071C | IMG2 | structural constituent of ribosome | 0.8113 | 1.8 | 9.61987E-05 |
| YDL098C | SNU23 | RNA splicing factor | 0.8125 | 1.8 | 0.0190395 |
| YHR214C-B | | | 0.8142 | 1.8 | 0.0902069 |
| YML045W | | | 0.8157 | 1.8 | 0.0319898 |
| YKL219W | COS9 | unknown | 0.8178 | 1.8 | 0.00531084 |
| YGR166W | KRE11 | unknown | 0.8188 | 1.8 | 0.0768148 |
| YDR410C | STE14 | protein-S-isoprenylcysteine O-methyltransferase | 0.8207 | 1.8 | 0.0164496 |
| YKL154W | SRP102 | GTPase * | 0.8232 | 1.8 | 0.0903301 |
| YFL045C | SEC53 | phosphomannomutase | 0.8275 | 1.8 | 0.00760547 |
| YGR021W | | unknown | 0.8277 | 1.8 | 0.0030717 |
| YNR077C | | | 0.8285 | 1.8 | 0.000991682 |
| YLR314C | CDC3 | structural constituent of cytoskeleton* | 0.8310 | 1.8 | 0.0150599 |
| YKL195W | MIA40 | unknown | 0.8312 | 1.8 | 0.0534365 |
| YER170W | ADK2 | adenylate kinase | 0.8363 | 1.8 | 0.0117105 |
| YIL117C | PRM5 | unknown | 0.8365 | 1.8 | 0.0331854 |
| YDR098C | GRX3 | thiol-disulfide exchange intermediate | 0.8378 | 1.8 | 0.358358 |
| YJL171C | | unknown | 0.8415 | 1.8 | 0.0114316 |
| YFL062W | COS4 | unknown | 0.8430 | 1.8 | 0.00660849 |
| YHR148W | IMP3 | snoRNA binding | 0.8437 | 1.8 | 0.0823706 |
| YMR067C | UBX4 | unknown | 0.8442 | 1.8 | 0.126909 |
| YGR295C | COS6 | unknown | 0.8463 | 1.8 | 0.000228968 |
| YDR244W | PEX5 | peroxisome targeting sequence binding | 0.8482 | 1.8 | 0.00416505 |
| YKL110C | KTI12 | enzyme regulator | 0.8497 | 1.8 | 0.00756375 |
| YLR146C | SPE4 | spermine synthase | 0.8508 | 1.8 | 0.00630244 |
| YBR160W | CDC28 | cyclin-dependent protein kinase | 0.8523 | 1.8 | 0.0435165 |
| YHR196W | UTP9 | snoRNA binding | 0.8540 | 1.8 | 0.156762 |
| YBR183W | YPC1 | ceramidase | 0.8553 | 1.8 | 0.000860177 |
| YGL127C | SOH1 | unknown | 0.8587 | 1.8 | 0.0410506 |
| YKL167C | MRP49 | structural constituent of ribosome | 0.8588 | 1.8 | 0.0167323 |
| YOL056W | GPM3 | unknown | 0.8605 | 1.8 | 0.111266 |
| YDL248W | COS7 | receptor | 0.8615 | 1.8 | 0.000649638 |
| YGL246C | RAI1 | enzyme regulator | 0.8630 | 1.8 | 0.0110987 |
| YNL037C | IDH1 | isocitrate dehydrogenase | 0.8665 | 1.8 | 0.0010171 |
| YCR003W | MRPL32 | structural constituent of ribosome | 0.8677 | 1.8 | 0.00542905 |
| YLR113W | HOG1 | MAP kinase | 0.8693 | 1.8 | 0.0263236 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-------------|
| YCR012W | PGK1 | phosphoglycerate kinase | 0.8720 | 1.8 | 0.0348544 |
| YBR088C | POL30 | DNA polymerase processivity factor | 0.8733 | 1.8 | 0.0399572 |
| YML132W | COS3 | protein binding | 0.8757 | 1.8 | 0.0164363 |
| YBR299W | MAL32 | alpha-glucosidase | 0.8757 | 1.8 | 0.0382171 |
| YJR057W | CDC8 | thymidylate kinase | 0.8765 | 1.8 | 0.0190524 |
| YFL-TYB | | | 0.8765 | 1.8 | 0.00677779 |
| YOL041C | NOP12 | RNA binding | 0.8768 | 1.8 | 0.0817458 |
| YDR181C | SAS4 | acetyltransferase * | 0.8770 | 1.8 | 0.027124 |
| YER187W | | unknown | 0.8777 | 1.8 | 0.0724041 |
| YNL193W | | unknown | 0.8780 | 1.8 | 0.0116237 |
| YKL087C | CYT2 | holocytochrome-c synthase | 0.8780 | 1.8 | 0.00591817 |
| YNL336W | COS1 | unknown | 0.8798 | 1.8 | 0.00122327 |
| YBR008C | FLR1 | multidrug transporter | 0.8800 | 1.8 | 0.0270774 |
| YFL063W | | | 0.8838 | 1.8 | 0.157521 |
| YKL060C | FBA1 | fructose-bisphosphate aldolase | 0.8840 | 1.8 | 0.0173647 |
| YHL048W | COS8 | unknown | 0.8908 | 1.9 | 0.00720741 |
| YPR060C | ARO7 | chorismate mutase | 0.8912 | 1.9 | 0.028491 |
| YOR292C | | unknown | 0.8977 | 1.9 | 0.0338396 |
| YGR207C | | unknown | 0.9017 | 1.9 | 0.00894998 |
| YNL334C | SNO2 | unknown | 0.9048 | 1.9 | 0.192218 |
| YDL202W | MRPL11 | structural constituent of ribosome | 0.9052 | 1.9 | 0.0193327 |
| YER053C | PIC2 | inorganic phosphate transporter | 0.9072 | 1.9 | 0.0398808 |
| YML025C | YML6 | structural constituent of ribosome | 0.9078 | 1.9 | 0.0275151 |
| YDR166C | SEC5 | protein binding | 0.9085 | 1.9 | 0.0844186 |
| YOR051C | | unknown | 0.9132 | 1.9 | 0.0737174 |
| YMR286W | MRPL33 | structural constituent of ribosome | 0.9163 | 1.9 | 0.0848865 |
| YMR304W | UBP15 | ubiquitin-specific protease | 0.9168 | 1.9 | 0.0045964 |
| YCL047C | | unknown | 0.9222 | 1.9 | 0.038372 |
| YOL122C | SMF1 | di-, tri-valent inorganic cation transporter | 0.9232 | 1.9 | 0.101084 |
| YBR014C | | unknown | 0.9267 | 1.9 | 0.0113188 |
| YPR138C | MEP3 | ammonium transporter | 0.9273 | 1.9 | 0.030441 |
| YOR021C | | unknown | 0.9305 | 1.9 | 0.000372097 |
| YKL001C | MET14 | adenylylsulfate kinase | 0.9312 | 1.9 | 0.0280655 |
| YLR332W | MID2 | transmembrane receptor | 0.9347 | 1.9 | 0.00696463 |
| YBR096W | | unknown | 0.9373 | 1.9 | 0.017836 |
| YGR091W | PRP31 | RNA splicing factor | 0.9398 | 1.9 | 0.000015137 |
| YKL067W | YNK1 | nucleoside diphosphate kinase | 0.9400 | 1.9 | 0.00192959 |
| YOL007C | | unknown | 0.9442 | 1.9 | 0.137754 |
| YIR043C | | unknown | 0.9458 | 1.9 | 0.00833612 |
| YNL030W | HHF2 | DNA binding | 0.9513 | 1.9 | 0.00389126 |
| YCL074W | | protein binding | 0.9518 | 1.9 | 0.012335 |
| YDR050C | TPI1 | triose-phosphate isomerase | 0.9577 | 1.9 | 0.032662 |
| YOL152W | FRE7 | ferric-chelate reductase | 0.9602 | 1.9 | 0.0174156 |
| YDL168W | SFA1 | alcohol dehydrogenase | 0.9602 | 1.9 | 0.0166407 |
| YBL028C | | unknown | 0.9605 | 1.9 | 0.00800421 |
| YKL096W | CWP1 | structural constituent of cell wall | 0.9623 | 1.9 | 0.00734584 |
| YKL066W | | | 0.9635 | 2.0 | 0.00245309 |
| YOR115C | TRS33 | unknown | 0.9653 | 2.0 | 0.0423615 |
| YLR376C | PSY3 | unknown | 0.9697 | 2.0 | 0.152877 |
| YDR322W | MRPL35 | structural constituent of ribosome | 0.9713 | 2.0 | 0.0111124 |
| YBL010C | | unknown | 0.9793 | 2.0 | 0.100959 |
| YLL009C | COX17 | thioredoxin peroxidase | 0.9812 | 2.0 | 0.0137056 |
| YGL012W | ERG4 | delta24(24-1) sterol reductase | 0.9812 | 2.0 | 0.113517 |
| YPL132W | COX11 | copper ion binding | 0.9823 | 2.0 | 0.00317913 |
| YHR144C | DCD1 | dCMP deaminase | 0.9898 | 2.0 | 0.0896276 |
| YPR158W | | unknown | 0.9975 | 2.0 | 0.0972752 |
| YHR173C | | | 0.9975 | 2.0 | 0.0185465 |
| YML027W | YOX1 | DNA binding | 0.9987 | 2.0 | 0.00759221 |
| YAL040C | CLN3 | cyclin-dependent protein kinase | 0.9995 | 2.0 | 0.00582507 |
| YHR178W | STB5 | transcription factor | 1.0032 | 2.0 | 0.00467186 |
| YHL004W | MRP4 | structural constituent of ribosome* | 1.0063 | 2.0 | 0.0104906 |
| YML087C | | unknown | 1.0160 | 2.0 | 0.0442256 |
| YBR282W | MRPL27 | structural constituent of ribosome | 1.0160 | 2.0 | 0.0598146 |
| YDL127W | PCL2 | cyclin-dependent protein kinase | 1.0177 | 2.0 | 0.00680612 |
| YOR131C | | unknown | 1.0182 | 2.0 | 0.0279522 |
| YNL300W | | unknown | 1.0182 | 2.0 | 0.0024862 |
| YKL200C | | | 1.0238 | 2.0 | 0.02866 |
| YMR225C | MRPL44 | structural constituent of ribosome | 1.0278 | 2.0 | 0.00105745 |
| YKR093W | PTR2 | peptide transporter | 1.0362 | 2.1 | 0.00845347 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|--|----------------------------|-------------|-------------|
| YPL223C | GRE1 | unknown | 1.0418 | 2.1 | 0.00294154 |
| YBL060W | | unknown | 1.0430 | 2.1 | 0.0213944 |
| YMR226C | TMA29 | oxidoreductase | 1.0443 | 2.1 | 0.0261447 |
| YDR511W | ACN9 | unknown | 1.0500 | 2.1 | 0.0208601 |
| YGL256W | ADH4 | alcohol dehydrogenase | 1.0537 | 2.1 | 0.00619427 |
| YNL231C | PDR16 | phosphatidylinositol transporter | 1.0583 | 2.1 | 0.014326 |
| YKL138C | MRPL31 | structural constituent of ribosome | 1.0590 | 2.1 | 0.0522296 |
| YGR215W | RSM27 | structural constituent of ribosome | 1.0600 | 2.1 | 0.0354891 |
| YIL060W | | unknown | 1.0655 | 2.1 | 0.00435071 |
| YHR059W | FYV4 | unknown | 1.0870 | 2.1 | 0.00671012 |
| YBL030C | PET9 | ATP:ADP antiporter | 1.0905 | 2.1 | 0.0378908 |
| YPR193C | HPA2 | histone acetyltransferase | 1.0942 | 2.1 | 0.00425003 |
| YMR193W | MRPL24 | structural constituent of ribosome | 1.0952 | 2.1 | 0.00351365 |
| YFR046C | CNN1 | unknown | 1.0982 | 2.1 | 0.0436128 |
| YNL122C | | unknown | 1.1042 | 2.1 | 0.00916503 |
| YDR333C | | unknown | 1.1045 | 2.2 | 0.00815121 |
| YFR035C | | unknown | 1.1080 | 2.2 | 0.0198341 |
| YKL113C | RAD27 | 5'-flap endonuclease | 1.1113 | 2.2 | 0.0294076 |
| YEL070W | DSF1 | unknown | 1.1157 | 2.2 | 0.0841826 |
| YEL040W | UTR2 | unknown | 1.1167 | 2.2 | 0.0947613 |
| YNL333W | SNZ2 | protein binding | 1.1173 | 2.2 | 0.0308249 |
| YMR190C | SGS1 | ATP-dependent DNA helicase | 1.1185 | 2.2 | 0.0439944 |
| YMR024W | MRPL3 | structural constituent of ribosome | 1.1188 | 2.2 | 0.0405321 |
| YNL213C | | unknown | 1.1202 | 2.2 | 0.00510172 |
| YDR115W | | structural constituent of ribosome | 1.1227 | 2.2 | 0.0181312 |
| YKR077W | | unknown | 1.1255 | 2.2 | 0.00211103 |
| YDL244W | THI13 | unknown | 1.1262 | 2.2 | 0.00209212 |
| YDL080C | THI3 | transcriptional activator * | 1.1267 | 2.2 | 0.0229008 |
| YHR145C | | | 1.1277 | 2.2 | 0.0262777 |
| YDL052C | SLC1 | 1-acylglycerol-3-phosphate O-acyltransferase | 1.1408 | 2.2 | 0.010186 |
| YER163C | | unknown | 1.1423 | 2.2 | 0.00683705 |
| YBR122C | MRPL36 | structural constituent of ribosome | 1.1518 | 2.2 | 0.00105675 |
| YGR084C | MRP13 | structural constituent of ribosome | 1.1530 | 2.2 | 0.0109299 |
| YHR046C | INM1 | inositol-1(or 4)-monophosphatase | 1.1563 | 2.2 | 0.000892312 |
| TDR261C-D | | | 1.1580 | 2.2 | 0.0213298 |
| YCL058C | | | 1.1597 | 2.2 | 0.0136091 |
| YMR316W | DIA1 | unknown | 1.1638 | 2.2 | 0.0186646 |
| YJR026W | | | 1.1652 | 2.2 | 0.0116285 |
| YMR199W | CLN1 | cyclin-dependent protein kinase | 1.1667 | 2.2 | 0.00162298 |
| YGR087C | PDC6 | pyruvate decarboxylase | 1.1667 | 2.2 | 0.00717395 |
| YER003C | PMI40 | mannose-6-phosphate isomerase | 1.1673 | 2.2 | 0.0020648 |
| YDR541C | | dihydrokaempferol 4-reductase | 1.1680 | 2.2 | 0.00599797 |
| YDR534C | FIT1 | unknown | 1.1717 | 2.3 | 0.00545346 |
| YML002W | | unknown | 1.1720 | 2.3 | 0.44113 |
| YKR006C | MRPL13 | structural constituent of ribosome | 1.1727 | 2.3 | 0.00361926 |
| YJR101W | RSM26 | structural constituent of ribosome | 1.1740 | 2.3 | 0.0029353 |
| YCR061W(1) | | | 1.1745 | 2.3 | 0.0373822 |
| YLR410W-B | | | 1.1752 | 2.3 | 0.00595837 |
| YKR016W | FMP13 | unknown | 1.1753 | 2.3 | 0.00356851 |
| YKL084W | HOT13 | unknown | 1.1792 | 2.3 | 0.0073707 |
| YBL059W | | unknown | 1.1900 | 2.3 | 0.0182627 |
| YKL137W | | unknown | 1.2010 | 2.3 | 0.0129598 |
| YNL155W | | unknown | 1.2022 | 2.3 | 0.0421847 |
| YBL101W-A | | | 1.2045 | 2.3 | 0.000528526 |
| YOR393W | ERR1 | phosphopyruvate hydratase | 1.2207 | 2.3 | 0.00967896 |
| YBR185C | MBA1 | unknown | 1.2248 | 2.3 | 0.00575463 |
| YOR286W | FMP31 | unknown | 1.2262 | 2.3 | 0.0170641 |
| YOR136W | IDH2 | isocitrate dehydrogenase | 1.2342 | 2.4 | 0.0133421 |
| YGL068W | MNP1 | unknown | 1.2360 | 2.4 | 0.0033026 |
| YBR106W | PHO88 | phosphate transporter | 1.2443 | 2.4 | 0.0176661 |
| YLR390W | ECM19 | unknown | 1.2453 | 2.4 | 0.0492809 |
| YLR044C | PDC1 | pyruvate decarboxylase | 1.2467 | 2.4 | 0.00115598 |
| YDR204W | COQ4 | unknown | 1.2493 | 2.4 | 0.00200241 |
| YDR399W | HPT1 | hypoxanthine phosphoribosyltransferase | 1.2495 | 2.4 | 0.0162737 |
| YMR184W | ADD37 | unknown | 1.2532 | 2.4 | 0.00862835 |
| YOR347C | PYK2 | pyruvate kinase | 1.2593 | 2.4 | 0.00107906 |
| YGR082W | TOM20 | protein transporter | 1.2612 | 2.4 | 0.00293878 |
| YDR316W-B | | | 1.2712 | 2.4 | 0.0025194 |
| YDR503C | LPP1 | phosphatidate phosphatase | 1.2727 | 2.4 | 0.0131322 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-------------|
| YAL038W | CDC19 | pyruvate kinase | 1.2733 | 2.4 | 0.00647856 |
| YMR002W | | unknown | 1.2852 | 2.4 | 0.00527914 |
| YFL059W | SNZ3 | protein binding | 1.2902 | 2.4 | 0.00473301 |
| YGR208W | SER2 | phosphoserine phosphatase | 1.2927 | 2.4 | 0.0112336 |
| YDR226W | ADK1 | adenylate kinase | 1.2943 | 2.5 | 0.00858258 |
| YIL044C | AGE2 | ARF GTPase activator | 1.2950 | 2.5 | 0.0391057 |
| YFL058W | THI5 | unknown | 1.3032 | 2.5 | 0.010547 |
| YJR029W | | | 1.3043 | 2.5 | 0.055112 |
| YGR222W | PET54 | RNA binding | 1.3100 | 2.5 | 0.0148642 |
| YHR174W | ENO2 | phosphopyruvate hydratase | 1.3123 | 2.5 | 0.00300659 |
| YHR038W | RRF1 | translation termination factor | 1.3212 | 2.5 | 0.0475878 |
| YHR087W | | unknown | 1.3237 | 2.5 | 0.00331972 |
| YNL070W | TOM7 | protein transporter | 1.3268 | 2.5 | 0.00272651 |
| YJR048W | CYC1 | electron carrier | 1.3350 | 2.5 | 0.00637233 |
| YCL020W | | | 1.3372 | 2.5 | 0.00301094 |
| YAL012W | CYS3 | cystathionine gamma-lyase | 1.3387 | 2.5 | 0.00509558 |
| YBR117C | TKL2 | transketolase | 1.3503 | 2.5 | 0.00690979 |
| YGL225W | VRG4 | nucleotide-sugar transporter | 1.3572 | 2.6 | 0.0110224 |
| YNL081C | SWS2 | structural constituent of ribosome | 1.3650 | 2.6 | 0.00855312 |
| YPL171C | OYE3 | NADPH dehydrogenase | 1.3667 | 2.6 | 0.010552 |
| YGR094W | VAS1 | valine-tRNA ligase | 1.3678 | 2.6 | 0.0163857 |
| YPL281C | ERR2 | phosphopyruvate hydratase | 1.3727 | 2.6 | 0.00511442 |
| YFL029C | CAK1 | cyclin-dependent protein kinase | 1.3787 | 2.6 | 0.00208134 |
| YAR030C | | | 1.3832 | 2.6 | 0.00504998 |
| YDR213W | UPC2 | RNA polymerase II transcription factor | 1.3865 | 2.6 | 0.00495435 |
| YCLX08C | | | 1.3932 | 2.6 | 0.0251234 |
| YNL185C | MRPL19 | structural constituent of ribosome | 1.4012 | 2.6 | 0.0124456 |
| YJR149W | | unknown | 1.4022 | 2.6 | 0.00746424 |
| YGL037C | PNC1 | nicotinamidase | 1.4173 | 2.7 | 0.000837184 |
| YEL039C | CYC7 | electron carrier | 1.4238 | 2.7 | 0.00310728 |
| YAR010C | | | 1.4320 | 2.7 | 0.00511234 |
| YHR181W | SVP26 | protein binding | 1.4367 | 2.7 | 0.0126539 |
| YBR012W-A | | | 1.4383 | 2.7 | 0.00946949 |
| YJL096W | MRPL49 | structural constituent of ribosome | 1.4423 | 2.7 | 0.000174505 |
| YER081W | SER3 | phosphoglycerate dehydrogenase | 1.4533 | 2.7 | 0.00242433 |
| YGR254W | ENO1 | phosphopyruvate hydratase | 1.4620 | 2.8 | 0.0153733 |
| YMR171C | | unknown | 1.4657 | 2.8 | 0.0804854 |
| YIL070C | MAM33 | unknown | 1.4880 | 2.8 | 0.00561867 |
| YLR227W-A | | | 1.4977 | 2.8 | 0.016713 |
| YFL051C | | unknown | 1.5142 | 2.9 | 0.0466355 |
| YOR385W | | unknown | 1.5202 | 2.9 | 0.00158224 |
| YHR104W | GRE3 | aldo-keto reductase * | 1.5240 | 2.9 | 0.00701331 |
| YER067W | | unknown | 1.5273 | 2.9 | 0.00258942 |
| YAL037C-B | | | 1.5282 | 2.9 | 0.00161913 |
| YCL075W | | protein binding | 1.5380 | 2.9 | 0.00508586 |
| YGL253W | HXK2 | hexokinase | 1.5510 | 2.9 | 0.00594349 |
| YHR179W | OYE2 | NADPH dehydrogenase | 1.5707 | 3.0 | 0.000618512 |
| YER062C | HOR2 | glycerol-1-phosphatase | 1.5950 | 3.0 | 0.00123764 |
| YGR142W | BTN2 | unknown | 1.5953 | 3.0 | 0.0357263 |
| YGR088W | CTT1 | catalase | 1.6050 | 3.0 | 0.000809239 |
| YIL053W | RHR2 | glycerol-1-phosphatase | 1.6108 | 3.1 | 0.00137762 |
| YBR116C | | | 1.6142 | 3.1 | 0.00182317 |
| YDL219W | DTD1 | hydrolase, acting on ester bonds | 1.6208 | 3.1 | 0.0135482 |
| YPL111W | CAR1 | zinc ion binding | 1.6462 | 3.1 | 0.0122879 |
| YDR391C | | unknown | 1.6593 | 3.2 | 0.00768137 |
| YGR151C | | | 1.7575 | 3.4 | 0.00548375 |
| YGL039W | | oxidoreductase | 1.7890 | 3.5 | 0.00140918 |
| YGL157W | | oxidoreductase | 1.8280 | 3.6 | 0.00128667 |
| YJR028W | | | 1.8443 | 3.6 | 0.00951414 |
| YFL021W | GAT1 | specific RNA polymerase II transcription factor | 1.8763 | 3.7 | 0.00226373 |
| YBL005W-A | | | 1.8923 | 3.7 | 0.00642121 |
| YDR400W | URH1 | ribosylpyrimidine nucleosidase | 1.8978 | 3.7 | 0.0295183 |
| YML040W | | | 1.9135 | 3.8 | 0.000847626 |
| YHR033W | | unknown | 1.9540 | 3.9 | 0.000486026 |
| YJL107C | | unknown | 2.0235 | 4.1 | 0.0361086 |
| YKL153W | | | 2.0405 | 4.1 | 0.00705793 |
| YFL013W-A | | | 2.2030 | 4.6 | 0.00213463 |
| YGR038C-A | | | 2.2115 | 4.6 | 0.00210448 |
| YFL014W | HSP12 | unknown | 2.2275 | 4.7 | 0.007048 |

Table 1: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--------------------------------------|----------------------------|-------------|-------------|
| YHR049W | FSH1 | serine hydrolase | 2.2737 | 4.8 | 0.00285462 |
| YLR035C-A | | | 2.2817 | 4.9 | 0.000973181 |
| YNL054W-B | | protein binding | 2.3353 | 5.0 | 0.00859456 |
| YPL163C | SVS1 | unknown | 2.3412 | 5.1 | 0.00346339 |
| YDR210C-D | | | 2.3965 | 5.3 | 0.00878016 |
| YKL152C | GPM1 | phosphoglycerate mutase | 2.4405 | 5.4 | 0.000740055 |
| YMR126C | DLT1 | unknown | 2.4488 | 5.5 | 0.00143872 |
| YLR168C | | unknown | 2.5583 | 5.9 | 0.00126568 |
| YOR382W | FIT2 | unknown | 2.6933 | 6.5 | 0.0105065 |
| YHR044C | DOG1 | 2-deoxyglucose-6-phosphatase | 2.8683 | 7.3 | 0.0112205 |
| YMR169C | ALD3 | 3-chloroallyl aldehyde dehydrogenase | 3.0673 | 8.4 | 0.00298248 |
| YJL108C | PRM10 | unknown | 3.0900 | 8.5 | 0.000916524 |
| YHR043C | DOG2 | 2-deoxyglucose-6-phosphatase | 3.1953 | 9.2 | 0.00846285 |
| YDR536W | STL1 | transporter | 3.5197 | 11.5 | 0.00666386 |

Table 2: Raw microarray data for fermentation day three.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|------------|---|----------------------------|-------------|-----------|
| YGR254W | ENO1 | phosphopyruvate hydratase | -3.6062 | -12.2 | 0.0015155 |
| YPL281C | ERR2 | phosphopyruvate hydratase | -3.2245 | -9.3 | 0.0036567 |
| YHR181W | SVP26 | protein binding | -3.1027 | -8.6 | 0.0019187 |
| YHR174W | ENO2 | phosphopyruvate hydratase | -3.1002 | -8.6 | 0.0036415 |
| YOR393W | ERR1 | phosphopyruvate hydratase | -2.9990 | -8.0 | 0.0340413 |
| YMR304C-A | | | -2.9637 | -7.8 | 0.0047968 |
| YCR012W | PGK1 | phosphoglycerate kinase | -2.8532 | -7.2 | 0.0138305 |
| YGR192C | TDH3 | glyceraldehyde-3-phosphate dehydrogenase | -2.7397 | -6.7 | 0.0002674 |
| YJL052W | TDH1 | glyceraldehyde-3-phosphate dehydrogenase | -2.7353 | -6.7 | 0.0021638 |
| YJR009C | TDH2 | glyceraldehyde-3-phosphate dehydrogenase | -2.5833 | -6.0 | 0.0004105 |
| YGR234W | YHB1 | nitric oxide reductase | -2.5823 | -6.0 | 0.0031394 |
| YGR240C | PFK1 | 6-phosphofructokinase | -2.5133 | -5.7 | 0.0125251 |
| YAL037C-B | | | -2.3255 | -5.0 | 0.0027095 |
| Q0250 | COX2 | cytochrome-c oxidase | -2.2987 | -4.9 | 0.0047764 |
| YLR044C | PDC1 | pyruvate decarboxylase | -2.2422 | -4.7 | 0.0099584 |
| YNL055C | POR1 | voltage-gated ion-selective channel | -2.1995 | -4.6 | 0.0098515 |
| YGR094W | VAS1 | valine-tRNA ligase | -2.1877 | -4.6 | 0.0015225 |
| YOL086C | ADH1 | alcohol dehydrogenase | -2.1872 | -4.6 | 0.013588 |
| YGR087C | PDC6 | pyruvate decarboxylase | -2.1603 | -4.5 | 1.429E-05 |
| YMR303C | ADH2 | alcohol dehydrogenase | -2.1495 | -4.4 | 0.0008692 |
| YLR134W | PDC5 | pyruvate decarboxylase | -2.1485 | -4.4 | 0.0264216 |
| YLR134W | PDC5 | pyruvate decarboxylase | -2.1485 | -4.4 | 0.0264216 |
| YDR050C | TPH1 | triose-phosphate isomerase | -2.1307 | -4.4 | 0.002872 |
| YJL005W | CYR1 | adenylate cyclase | -2.1205 | -4.3 | 0.0104521 |
| YBR145W | ADH5 | alcohol dehydrogenase | -2.0963 | -4.3 | 0.0013869 |
| YPL187W | MF(ALPHA)1 | mating pheromone | -2.0948 | -4.3 | 0.0110709 |
| YHR173C | | | -2.0733 | -4.2 | 0.0204469 |
| YBR144C | | | -2.0662 | -4.2 | 0.0025647 |
| YNL015W | PBI2 | endopeptidase inhibitor | -2.0497 | -4.1 | 0.0064836 |
| YNL072W | RNH201 | ribonuclease H | -2.0482 | -4.1 | 0.0054351 |
| YKL152C | GPM1 | phosphoglycerate mutase | -2.0472 | -4.1 | 0.0012347 |
| YGL089C | MF(ALPHA)2 | mating pheromone | -2.0032 | -4.0 | 0.0023197 |
| YKL060C | FBA1 | fructose-bisphosphate aldolase | -1.9920 | -4.0 | 0.0020741 |
| YKL174C | TPO5 | polyamine transporter | -1.9703 | -3.9 | 0.0053626 |
| YFL030W | AGX1 | alanine-glyoxylate transaminase | -1.9673 | -3.9 | 0.0015463 |
| YFL011W | HXT10 | glucose transporter | -1.9588 | -3.9 | 0.0058639 |
| YLR153C | ACS2 | acetate-CoA ligase | -1.8628 | -3.6 | 0.0009635 |
| YPR160W | GPH1 | glycogen phosphorylase | -1.8208 | -3.5 | 0.0123043 |
| YPR160W | GPH1 | glycogen phosphorylase | -1.8208 | -3.5 | 0.0123043 |
| YAL038W | CDC19 | pyruvate kinase | -1.8033 | -3.5 | 0.0001967 |
| YDR246W | TRS23 | unknown | -1.7960 | -3.5 | 0.0019749 |
| YBL030C | PET9 | ATP:ADP antiporter | -1.7840 | -3.4 | 0.0121792 |
| YJR121W | ATP2 | hydrogen-transporting ATP synthase , rotational mechanism | -1.7665 | -3.4 | 0.0013171 |
| YPL262W | FUM1 | fumarate hydratase | -1.7648 | -3.4 | 0.0080621 |
| YFL018C | LPD1 | glycine dehydrogenase | -1.7585 | -3.4 | 0.0113803 |
| YBL099W | ATP1 | hydrogen-transporting ATP synthase , rotational mechanism | -1.7580 | -3.4 | 0.0004718 |
| YNL135C | FPR1 | peptidyl-prolyl cis-trans isomerase | -1.7398 | -3.3 | 0.0023218 |
| YLR259C | HSP60 | single-stranded DNA binding | -1.7267 | -3.3 | 0.0154674 |
| YJR048W | CYC1 | electron carrier | -1.7030 | -3.3 | 0.0050096 |
| YKL153W | | | -1.6862 | -3.2 | 0.0036681 |
| YMR083W | ADH3 | alcohol dehydrogenase | -1.6602 | -3.2 | 0.017519 |
| YDR247W | VHS1 | protein kinase | -1.6598 | -3.2 | 0.0099291 |
| YDR249C | | unknown | -1.6517 | -3.1 | 0.0097933 |
| YEL039C | CYC7 | electron carrier | -1.6505 | -3.1 | 0.0038712 |
| YDR453C | TSA2 | thioredoxin peroxidase | -1.5847 | -3.0 | 0.017825 |
| YJR016C | ILV3 | dihydroxy-acid dehydratase | -1.5752 | -3.0 | 0.0011641 |
| YBR085W | AAC3 | ATP:ADP antiporter | -1.5737 | -3.0 | 0.0001426 |
| YGL179C | TOS3 | protein kinase | -1.5720 | -3.0 | 0.004032 |
| YHR092C | HXT4 | glucose transporter | -1.5598 | -2.9 | 0.0322082 |
| YBR078W | ECM33 | unknown | -1.5307 | -2.9 | 0.0004679 |
| YER133W | GLC7 | protein phosphatase type 1 | -1.5160 | -2.9 | 0.0195183 |
| YML028W | TSA1 | thioredoxin peroxidase | -1.5123 | -2.9 | 0.0050001 |
| YNL053W | MSG5 | prenylated protein tyrosine phosphatase | -1.5088 | -2.8 | 6.246E-05 |
| YDR380W | ARO10 | pyruvate decarboxylase | -1.4992 | -2.8 | 0.007323 |
| YMR190C | SGS1 | ATP-dependent DNA helicase | -1.4945 | -2.8 | 0.047043 |
| YGR244C | LSC2 | succinate-CoA ligase | -1.4912 | -2.8 | 0.0171706 |
| YKL148C | SDH1 | succinate dehydrogenase | -1.4842 | -2.8 | 0.0233448 |
| YBL001C | ECM15 | unknown | -1.4735 | -2.8 | 0.0042214 |
| YNL052W | COX5A | cytochrome-c oxidase | -1.4733 | -2.8 | 0.0017709 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YCR074C | | | -1.4615 | -2.8 | 0.0186611 |
| YHR051W | COX6 | cytochrome-c oxidase | -1.4555 | -2.7 | 0.0053019 |
| YJL153C | INO1 | inositol-3-phosphate synthase | -1.4540 | -2.7 | 0.0129276 |
| YLR395C | COX8 | cytochrome-c oxidase | -1.4505 | -2.7 | 0.0040148 |
| YER023W | PRO3 | pyrroline-5-carboxylate reductase | -1.4472 | -2.7 | 0.0083757 |
| YCR013C | | | -1.4433 | -2.7 | 0.178912 |
| YHR008C | SOD2 | manganese superoxide dismutase | -1.4285 | -2.7 | 0.001679 |
| YGL191W | COX13 | enzyme regulator * | -1.4207 | -2.7 | 0.0001805 |
| YLL041C | SDH2 | succinate dehydrogenase | -1.4120 | -2.7 | 5.206E-05 |
| YNL103W | MET4 | transcription coactivator | -1.4093 | -2.7 | 0.0002534 |
| YGR279C | SCW4 | glucosidase | -1.4013 | -2.6 | 0.0438594 |
| YJL152W | | | -1.3928 | -2.6 | 0.0041552 |
| YOR215C | | unknown | -1.3892 | -2.6 | 0.0227276 |
| YHR001W-A | QCR10 | ubiquinol-cytochrome-c reductase | -1.3537 | -2.6 | 0.0086701 |
| YDR322W | MRPL35 | structural constituent of ribosome | -1.3517 | -2.6 | 0.000573 |
| YGR246C | BRF1 | RNA polymerase III transcription factor | -1.3508 | -2.6 | 0.0326688 |
| YKL141W | SDH3 | succinate dehydrogenase | -1.3158 | -2.5 | 0.0037429 |
| YBL100C | | | -1.3097 | -2.5 | 0.0323298 |
| YDR273W | DON1 | unknown | -1.2945 | -2.5 | 0.0016658 |
| YIL155C | GUT2 | glycerol-3-phosphate dehydrogenase | -1.2882 | -2.4 | 0.0274819 |
| YDL067C | COX9 | cytochrome-c oxidase | -1.2878 | -2.4 | 0.0461493 |
| YKL160W | ELF1 | RNA polymerase II transcription elongation factor | -1.2837 | -2.4 | 0.0017938 |
| YJR120W | | unknown | -1.2828 | -2.4 | 0.0003826 |
| YDR119W | | unknown | -1.2732 | -2.4 | 0.0078908 |
| YHR219W | | unknown | -1.2722 | -2.4 | 0.0171083 |
| YBL098W | BNA4 | kynurenine 3-monooxygenase | -1.2688 | -2.4 | 0.0055453 |
| YKL085W | MDH1 | L-malate dehydrogenase | -1.2673 | -2.4 | 0.0038412 |
| YJL217W | | unknown | -1.2667 | -2.4 | 0.0479643 |
| YJL217W | | unknown | -1.2667 | -2.4 | 0.0479643 |
| YHR094C | HXT1 | glucose transporter | -1.2655 | -2.4 | 0.0098239 |
| YJL214W | HXT8 | glucose transporter | -1.2638 | -2.4 | 0.0358475 |
| YJL066C | MPM1 | unknown | -1.2535 | -2.4 | 0.015919 |
| YJR148W | BAT2 | branched-chain-amino-acid transaminase | -1.2505 | -2.4 | 0.007391 |
| YDR245W | MNN10 | alpha-1,6-mannosyltransferase | -1.2335 | -2.4 | 0.0113862 |
| YDR202C | RAV2 | unknown | -1.2270 | -2.3 | 0.0135408 |
| YOR133W | EFT1 | translation elongation factor | -1.2250 | -2.3 | 0.0679752 |
| YMR145C | NDE1 | NADH dehydrogenase | -1.2160 | -2.3 | 0.0949257 |
| YKL067W | YNK1 | nucleoside diphosphate kinase | -1.2115 | -2.3 | 0.0016627 |
| YBL015W | ACH1 | acetyl-CoA hydrolase | -1.2090 | -2.3 | 0.0080694 |
| YKL164C | PIR1 | structural constituent of cell wall | -1.2010 | -2.3 | 0.0299198 |
| YJL045W | | succinate dehydrogenase | -1.1980 | -2.3 | 0.0009649 |
| YKL145W | RPT1 | ATPase * | -1.1978 | -2.3 | 0.0374322 |
| YKL145W | RPT1 | ATPase * | -1.1978 | -2.3 | 0.0374322 |
| YKL026C | GPX1 | glutathione peroxidase * | -1.1797 | -2.3 | 0.0193568 |
| YOR230W | WTM1 | transcription corepressor | -1.1757 | -2.3 | 0.0049473 |
| YKL066W | | | -1.1743 | -2.3 | 0.0367139 |
| YER178W | PDA1 | pyruvate dehydrogenase | -1.1707 | -2.3 | 0.0048228 |
| YMR256C | COX7 | cytochrome-c oxidase | -1.1678 | -2.2 | 0.002045 |
| YIL170W | HXT12 | unknown | -1.1670 | -2.2 | 0.0020793 |
| YIL170W | HXT12 | unknown | -1.1670 | -2.2 | 0.0020793 |
| YGL253W | HXK2 | hexokinase | -1.1618 | -2.2 | 0.0208222 |
| YEL030W | ECM10 | unknown | -1.1598 | -2.2 | 0.13813 |
| YJR077C | MIR1 | inorganic phosphate transporter | -1.1593 | -2.2 | 0.0012406 |
| YLL066W-A | | | -1.1532 | -2.2 | 0.0420793 |
| YPL134C | ODC1 | intracellular transporter | -1.1513 | -2.2 | 0.005421 |
| YHR007C | ERG11 | sterol 14-demethylase | -1.1313 | -2.2 | 0.0215179 |
| YNL220W | ADE12 | adenylosuccinate synthase | -1.1282 | -2.2 | 0.0175976 |
| YFL038C | YPT1 | GTPase | -1.1268 | -2.2 | 0.008804 |
| YBR039W | ATP3 | hydrogen-transporting ATP synthase , rotational mechanism | -1.1240 | -2.2 | 0.0552641 |
| YIR037W | HYR1 | thioredoxin peroxidase * | -1.1185 | -2.2 | 0.0279486 |
| YBR294W | SUL1 | sulfate transporter | -1.1145 | -2.2 | 0.0080761 |
| YDR385W | EFT2 | translation elongation factor | -1.1065 | -2.2 | 0.0312713 |
| YAR018C | KIN3 | protein kinase | -1.1060 | -2.2 | 0.0051538 |
| YGL105W | ARC1 | tRNA binding | -1.1048 | -2.2 | 0.0153682 |
| YLR303W | MET17 | cysteine synthase * | -1.1033 | -2.1 | 0.0546932 |
| YBR244W | GPX2 | glutathione peroxidase * | -1.1027 | -2.1 | 0.0007172 |
| YBR093C | PHO5 | acid phosphatase * | -1.1022 | -2.1 | 0.0198011 |
| YCL058C | | | -1.1002 | -2.1 | 0.0466576 |
| YJR073C | OPI3 | phosphatidyl-N-methylethanolamine N-methyltransferase | -1.0943 | -2.1 | 0.0047786 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YEL024W | RIP1 | ubiquinol-cytochrome-c reductase | -1.0880 | -2.1 | 0.0069601 |
| YLR294C | | | -1.0695 | -2.1 | 0.0021397 |
| YEL066W | HPA3 | histone acetyltransferase | -1.0658 | -2.1 | 0.0229021 |
| YEL066W | HPA3 | histone acetyltransferase | -1.0658 | -2.1 | 0.0229021 |
| YFL039C | ACT1 | structural constituent of cytoskeleton | -1.0655 | -2.1 | 0.0120042 |
| YKL016C | ATP7 | structural molecule * | -1.0593 | -2.1 | 0.0016204 |
| YDR320C | SWA2 | protein binding | -1.0588 | -2.1 | 0.0431103 |
| YHL050C | | helicase | -1.0503 | -2.1 | 0.0031794 |
| YKL097W-A | | | -1.0437 | -2.1 | 0.0006083 |
| YML110C | COQ5 | 2-hexaprenyl-6-methoxy-1,4-benzoquinone methyltransferase | -1.0390 | -2.1 | 0.0479456 |
| YHR214W | | unknown | -1.0318 | -2.0 | 0.0180394 |
| YPR020W | ATP20 | structural molecule | -1.0255 | -2.0 | 0.0010649 |
| YEL077C | | helicase | -1.0240 | -2.0 | 0.002784 |
| YEL065W | SIT1 | siderophore-iron (ferrioxamine) uptake transporter | -1.0200 | -2.0 | 0.0134719 |
| YBL113W-A | | | -1.0152 | -2.0 | 0.0042373 |
| YHL034C | SBP1 | RNA binding | -1.0132 | -2.0 | 0.0048516 |
| YPL078C | ATP4 | structural molecule * | -1.0002 | -2.0 | 0.0020795 |
| YDL181W | INH1 | enzyme inhibitor | -0.9997 | -2.0 | 0.0071249 |
| YCR004C | YCP4 | electron carrier | -0.9735 | -2.0 | 0.014026 |
| YLR320W | MMS22 | unknown | -0.9560 | -1.9 | 0.0562386 |
| YDR438W | | unknown | -0.9538 | -1.9 | 0.0397923 |
| YHR126C | | unknown | -0.9492 | -1.9 | 0.0112156 |
| YDL055C | PSA1 | mannose-1-phosphate guanylyltransferase | -0.9452 | -1.9 | 0.0086477 |
| YBL002W | HTB2 | DNA binding | -0.9392 | -1.9 | 0.0188168 |
| YJL143W | TIM17 | protein transporter | -0.9383 | -1.9 | 0.0650109 |
| YLR150W | STM1 | DNA binding* | -0.9362 | -1.9 | 0.0100821 |
| YBR067C | TIP1 | structural constituent of cell wall* | -0.9323 | -1.9 | 0.0027268 |
| YDR268W | MSW1 | tryptophan-tRNA ligase | -0.9292 | -1.9 | 0.0036085 |
| YDL078C | MDH3 | L-malate dehydrogenase | -0.9092 | -1.9 | 0.0225068 |
| YNL037C | IDH1 | isocitrate dehydrogenase | -0.9092 | -1.9 | 0.0060415 |
| YGR182C | | | -0.9003 | -1.9 | 0.0346738 |
| YAL008W | FUN14 | unknown | -0.9000 | -1.9 | 0.109014 |
| YFR003C | YPII | protein phosphatase inhibitor | -0.8968 | -1.9 | 6.043E-05 |
| YDL137W | ARF2 | GTPase | -0.8950 | -1.9 | 0.0160564 |
| YER011W | TIR1 | structural constituent of cell wall | -0.8910 | -1.9 | 0.0206417 |
| YDL215C | GDH2 | glutamate dehydrogenase | -0.8852 | -1.8 | 0.0142824 |
| YNL064C | YDJ1 | chaperone regulator * | -0.8830 | -1.8 | 0.0830227 |
| YKR013W | PRY2 | unknown | -0.8778 | -1.8 | 0.0225798 |
| YBL078C | ATG8 | microtubule binding | -0.8757 | -1.8 | 0.0573605 |
| YDL021W | GPM2 | unknown | -0.8677 | -1.8 | 0.0045561 |
| YDL021W | GPM2 | unknown | -0.8677 | -1.8 | 0.0045561 |
| YML132W | COS3 | protein binding | -0.8665 | -1.8 | 0.0077908 |
| YDR529C | QCR7 | ubiquinol-cytochrome-c reductase | -0.8472 | -1.8 | 0.0060546 |
| YBR025C | | unknown | -0.8447 | -1.8 | 0.0049425 |
| YJR103W | URA8 | CTP synthase | -0.8405 | -1.8 | 0.0835693 |
| YDR193W | | | -0.8392 | -1.8 | 0.0085795 |
| YDR033W | MRH1 | unknown | -0.8385 | -1.8 | 0.0005238 |
| YLR377C | FBP1 | fructose-bisphosphatase | -0.8233 | -1.8 | 0.0738824 |
| YNL142W | MEP2 | ammonium transporter | -0.8227 | -1.8 | 0.111587 |
| YFR053C | HXK1 | hexokinase | -0.8142 | -1.8 | 0.0146429 |
| YLL050C | COF1 | protein binding | -0.8098 | -1.8 | 0.101651 |
| YIL111W | COX5B | cytochrome-c oxidase | -0.8067 | -1.7 | 0.0204686 |
| YKL117W | SBA1 | chaperone binding | -0.7973 | -1.7 | 0.0162102 |
| YER053C | PIC2 | inorganic phosphate transporter | -0.7935 | -1.7 | 0.004651 |
| YNR001C | CIT1 | citrate (Si)-synthase | -0.7907 | -1.7 | 0.0039411 |
| YEL009C | GCN4 | DNA binding* | -0.7893 | -1.7 | 0.0013036 |
| YLR258W | GSY2 | glycogen (starch) synthase | -0.7845 | -1.7 | 0.131435 |
| YOR027W | STI1 | unfolded protein binding* | -0.7800 | -1.7 | 0.103367 |
| YPL037C | EGD1 | unfolded protein binding | -0.7800 | -1.7 | 0.0296889 |
| YBR094W | | unknown | -0.7758 | -1.7 | 0.0068261 |
| YDL192W | ARF1 | GTPase | -0.7753 | -1.7 | 0.0196191 |
| TMR307W | | | -0.7728 | -1.7 | 0.0020019 |
| YPL271W | ATP15 | hydrogen-transporting ATP synthase , rotational mechanism | -0.7705 | -1.7 | 0.0419803 |
| YJL001W | PRE3 | endopeptidase | -0.7700 | -1.7 | 0.0661903 |
| YIL123W | SIM1 | unknown | -0.7668 | -1.7 | 0.0555948 |
| YKL192C | ACP1 | acyl carrier | -0.7663 | -1.7 | 0.0116203 |
| YFR049W | YMR31 | structural constituent of ribosome | -0.7653 | -1.7 | 0.0024639 |
| YFR049W | YMR31 | structural constituent of ribosome | -0.7653 | -1.7 | 0.0024639 |
| YJR031C | GEA1 | ARF guanyl-nucleotide exchange factor | -0.7608 | -1.7 | 0.0114537 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YLL026W | HSP104 | chaperone binding* | -0.7602 | -1.7 | 0.0250434 |
| YHR040W | BCD1 | nucleic acid binding | -0.7593 | -1.7 | 0.0289917 |
| YFL062W | COS4 | unknown | -0.7565 | -1.7 | 0.0238822 |
| YBL075C | SSA3 | ATPase * | -0.7562 | -1.7 | 0.0299285 |
| YBR054W | YRO2 | unknown | -0.7555 | -1.7 | 0.024774 |
| YIL166C | | transporter | -0.7520 | -1.7 | 0.0242363 |
| YJR045C | SSC1 | ATPase * | -0.7508 | -1.7 | 0.0233663 |
| YBL045C | COR1 | ubiquinol-cytochrome-c reductase | -0.7493 | -1.7 | 0.0293268 |
| YJL219W | HXT9 | glucose transporter | -0.7388 | -1.7 | 0.152974 |
| YDR284C | DPP1 | phosphatidate phosphatase * | -0.7358 | -1.7 | 0.0849586 |
| YDR032C | PST2 | unknown | -0.7285 | -1.7 | 0.0423654 |
| YLR304C | ACO1 | aconitate hydratase | -0.7272 | -1.7 | 0.0083209 |
| YOR136W | IDH2 | isocitrate dehydrogenase | -0.7252 | -1.7 | 0.0740756 |
| YDL066W | IDP1 | isocitrate dehydrogenase | -0.7230 | -1.7 | 0.0028387 |
| YJR001W | AVT1 | neutral amino acid transporter | -0.7193 | -1.6 | 0.0043982 |
| YJL068C | | S-formylglutathione hydrolase | -0.7135 | -1.6 | 0.0044765 |
| YDR133C | | | -0.7077 | -1.6 | 0.0340978 |
| YDR134C | | unknown | -0.7057 | -1.6 | 0.0016352 |
| YGL062W | PYC1 | pyruvate carboxylase | -0.7055 | -1.6 | 0.031003 |
| YIL136W | OM45 | unknown | -0.7037 | -1.6 | 0.0417209 |
| YDL029W | ARP2 | structural constituent of cytoskeleton* | -0.6987 | -1.6 | 0.0857069 |
| YMR203W | TOM40 | protein transporter | -0.6987 | -1.6 | 0.000612 |
| YNL009W | IDP3 | isocitrate dehydrogenase | -0.6985 | -1.6 | 0.003158 |
| YJL034W | KAR2 | ATPase * | -0.6963 | -1.6 | 0.0618993 |
| YJR008W | | unknown | -0.6962 | -1.6 | 0.0435149 |
| YJR008W | | unknown | -0.6962 | -1.6 | 0.0435149 |
| YJL200C | ACO2 | aconitate hydratase | -0.6947 | -1.6 | 0.096495 |
| YGL200C | EMP24 | unknown | -0.6932 | -1.6 | 0.04933 |
| YEL034W | HYP2 | protein binding* | -0.6910 | -1.6 | 0.0130075 |
| YDR327W | | | -0.6852 | -1.6 | 0.0101475 |
| YDR327W | | | -0.6852 | -1.6 | 0.0101475 |
| YBR139W | | serine hydrolase * | -0.6830 | -1.6 | 0.0061513 |
| YMR295C | | unknown | -0.6830 | -1.6 | 0.0392713 |
| YNL007C | SIS1 | unfolded protein binding* | -0.6815 | -1.6 | 0.0014455 |
| YDL100C | GET3 | ATPase | -0.6792 | -1.6 | 0.0208688 |
| YMR297W | PRC1 | carboxypeptidase C | -0.6787 | -1.6 | 0.0625163 |
| YML078W | CPR3 | peptidyl-prolyl cis-trans isomerase | -0.6785 | -1.6 | 0.006109 |
| YGL121C | GPG1 | signal transducer | -0.6768 | -1.6 | 0.0036963 |
| YCL043C | PDH1 | protein disulfide isomerase | -0.6720 | -1.6 | 0.0105996 |
| YIL138C | TPM2 | actin lateral binding | -0.6720 | -1.6 | 0.0003805 |
| YBR080C | SEC18 | ATPase | -0.6645 | -1.6 | 0.0970178 |
| YDL206W | | unknown | -0.6642 | -1.6 | 0.0344768 |
| YOR020C | HSP10 | unfolded protein binding | -0.6618 | -1.6 | 0.032411 |
| YER004W | FMP52 | unknown | -0.6575 | -1.6 | 0.0297763 |
| YDR382W | RPP2B | structural constituent of ribosome | -0.6533 | -1.6 | 0.0767667 |
| YHL034W-A | | | -0.6515 | -1.6 | 0.0723914 |
| YIL033C | BCY1 | cAMP-dependent protein kinase | -0.6437 | -1.6 | 0.0054367 |
| YDR377W | ATP17 | hydrogen-transporting ATP synthase , rotational mechanism | -0.6413 | -1.6 | 0.0019884 |
| YDR148C | KGD2 | dihydrolipoyllysine-residue succinyltransferase | -0.6405 | -1.6 | 0.0036853 |
| YBR221C | PDB1 | pyruvate dehydrogenase | -0.6392 | -1.6 | 0.0672441 |
| YHR215W | PHO12 | acid phosphatase | -0.6392 | -1.6 | 0.0412283 |
| YMR260C | TIF11 | translation initiation factor | -0.6378 | -1.6 | 0.0377335 |
| YLR248W | RCK2 | protein serine/threonine kinase | -0.6287 | -1.5 | 0.0998243 |
| YEL034C-A | | | -0.6280 | -1.5 | 0.0763997 |
| YJL158C | CIS3 | structural constituent of cell wall | -0.6232 | -1.5 | 0.0386436 |
| YDL126C | CDC48 | ATPase | -0.6207 | -1.5 | 0.125145 |
| YDR028C | REG1 | protein phosphatase type 1 regulator | -0.6190 | -1.5 | 0.160949 |
| YOR007C | SGT2 | unknown | -0.6023 | -1.5 | 0.0440004 |
| YIL018W | RPL2B | structural constituent of ribosome | -0.6010 | -1.5 | 0.0942016 |
| YAL013W | DEP1 | transcription regulator | -0.6008 | -1.5 | 0.0155276 |
| YLR149C | | unknown | -0.6003 | -1.5 | 0.0570284 |
| YLR043C | TRX1 | thiol-disulfide exchange intermediate | -0.6002 | -1.5 | 0.0166047 |
| YBR243C | ALG7 | UDP-N-acetylglucosamine-dolichyl-phosphate N-acetylglucosaminyl transferase | -0.6000 | -1.5 | 0.0112551 |
| YOL030W | GAS5 | 1,3-beta-glucanosyltransferase | -0.5978 | -1.5 | 0.0491241 |
| YJL026W | RNR2 | ribonucleoside-diphosphate reductase | -0.5912 | -1.5 | 0.133042 |
| YCL035C | GRX1 | thiol-disulfide exchange intermediate * | -0.5895 | -1.5 | 0.0611118 |
| YPR035W | GLN1 | glutamate-ammonia ligase | -0.5880 | -1.5 | 0.0211799 |
| YER094C | PUP3 | endopeptidase | -0.5863 | -1.5 | 0.0298535 |
| YML106W | URA5 | orotate phosphoribosyltransferase | -0.5833 | -1.5 | 0.0639159 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YGR253C | PUP2 | endopeptidase | -0.5820 | -1.5 | 0.152895 |
| YHR214W-A | | | -0.5805 | -1.5 | 0.0033847 |
| YHR057C | CPR2 | peptidyl-prolyl cis-trans isomerase | -0.5798 | -1.5 | 5.575E-05 |
| YHR057C | CPR2 | peptidyl-prolyl cis-trans isomerase | -0.5798 | -1.5 | 5.575E-05 |
| YCL009C | ILV6 | enzyme regulator * | -0.5775 | -1.5 | 0.147573 |
| YIL011W | TIR3 | unknown | -0.5722 | -1.5 | 0.0526976 |
| YPL033C | | unknown | -0.5700 | -1.5 | 0.0180346 |
| YPL033C | | unknown | -0.5700 | -1.5 | 0.0180346 |
| YKL216W | URA1 | dihydroorotate dehydrogenase | -0.5687 | -1.5 | 0.0004113 |
| YER086W | ILV1 | threonine ammonia-lyase | -0.5685 | -1.5 | 0.214924 |
| YHL048W | COS8 | unknown | -0.5677 | -1.5 | 0.0700487 |
| YBR164C | ARL1 | GTPase | -0.5667 | -1.5 | 0.0294648 |
| YNR036C | | structural constituent of ribosome | -0.5595 | -1.5 | 0.0608814 |
| YDR086C | SSS1 | protein transporter | -0.5593 | -1.5 | 0.0117785 |
| YBL003C | HTA2 | DNA binding | -0.5588 | -1.5 | 0.0139608 |
| YJL159W | HSP150 | structural constituent of cell wall | -0.5577 | -1.5 | 0.0256999 |
| YDL007W | RPT2 | ATPase * | -0.5568 | -1.5 | 0.0535129 |
| YGR183C | QCR9 | ubiquinol-cytochrome-c reductase | -0.5528 | -1.5 | 0.0528061 |
| YHR192W | | unknown | -0.5472 | -1.5 | 0.0007977 |
| YIR041W | PAU15 | unknown | -0.5462 | -1.5 | 0.053258 |
| YGR271W | SLH1 | RNA helicase | -0.5458 | -1.5 | 0.058332 |
| YDR321W | ASP1 | asparaginase | -0.5450 | -1.5 | 0.0152668 |
| YPL270W | MDL2 | ATPase , coupled to transmembrane movement of substances | -0.5438 | -1.5 | 0.0662755 |
| YNL079C | TPM1 | actin lateral binding | -0.5402 | -1.5 | 0.185009 |
| YJL207C | LAA1 | unknown | -0.5393 | -1.5 | 0.0113212 |
| YBR111C | YSA1 | phosphoribosyl-ATP diphosphatase | -0.5387 | -1.5 | 0.283746 |
| YMR056C | AAC1 | ATP:ADP antiporter | -0.5370 | -1.5 | 0.0662925 |
| YDR163W | CWC15 | unknown | -0.5333 | -1.4 | 0.0393203 |
| YGR282C | BGL2 | glucan 1,3-beta-glucosidase | -0.5302 | -1.4 | 0.0013714 |
| YLR174W | IDP2 | isocitrate dehydrogenase | -0.5297 | -1.4 | 0.0493384 |
| YDR388W | RVS167 | cytoskeletal protein binding | -0.5288 | -1.4 | 0.0285552 |
| YNL208W | | unknown | -0.5277 | -1.4 | 0.109385 |
| YAR020C | PAU7 | unknown | -0.5165 | -1.4 | 8.053E-05 |
| YBR148W | YSW1 | unknown | -0.5153 | -1.4 | 0.001374 |
| YMR205C | PFK2 | 6-phosphofructokinase | -0.5130 | -1.4 | 0.142388 |
| YBR184W | | unknown | -0.5118 | -1.4 | 0.308085 |
| YGR180C | RNR4 | ribonucleoside-diphosphate reductase | -0.5117 | -1.4 | 0.0193124 |
| YER177W | BMH1 | protein binding* | -0.5098 | -1.4 | 0.141298 |
| YLR257W | | unknown | -0.5087 | -1.4 | 0.0086452 |
| YEL032W | MCM3 | chromatin binding* | -0.5028 | -1.4 | 0.0398392 |
| YGR209C | TRX2 | thiol-disulfide exchange intermediate | -0.4998 | -1.4 | 0.0114102 |
| YJL167W | ERG20 | dimethylallyltranstransferase * | -0.4997 | -1.4 | 0.0353727 |
| YDR198C | RKM2 | unknown | -0.4978 | -1.4 | 0.0080298 |
| YDR198C | RKM2 | unknown | -0.4978 | -1.4 | 0.0080298 |
| YAL023C | PMT2 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.4962 | -1.4 | 0.0299618 |
| YIL176C | PAU14 | unknown | -0.4953 | -1.4 | 0.0086783 |
| YBR023C | CHS3 | chitin synthase | -0.4952 | -1.4 | 0.0954594 |
| YLL004W | ORC3 | DNA replication origin binding | -0.4935 | -1.4 | 0.0167494 |
| YLL004W | ORC3 | DNA replication origin binding | -0.4935 | -1.4 | 0.0167494 |
| YLR229C | CDC42 | GTPase * | -0.4898 | -1.4 | 0.0031201 |
| YAL005C | SSA1 | ATPase * | -0.4895 | -1.4 | 0.0431829 |
| YEL025C | | unknown | -0.4887 | -1.4 | 0.218472 |
| YIL125W | KGD1 | oxoglutarate dehydrogenase | -0.4863 | -1.4 | 0.037867 |
| YDR454C | GUK1 | guanylate kinase | -0.4852 | -1.4 | 0.0938369 |
| YBR072W | HSP26 | unfolded protein binding | -0.4848 | -1.4 | 0.0065322 |
| YDL130W | RPP1B | structural constituent of ribosome | -0.4847 | -1.4 | 0.0816368 |
| YJR047C | ANB1 | translation initiation factor | -0.4833 | -1.4 | 0.010185 |
| YAL063C | FLO9 | mannose binding | -0.4823 | -1.4 | 0.0359755 |
| YLR110C | CCW12 | unknown | -0.4818 | -1.4 | 0.0774241 |
| YNL124W | NAF1 | RNA binding | -0.4817 | -1.4 | 0.218076 |
| YOR362C | PRE10 | endopeptidase | -0.4783 | -1.4 | 0.258326 |
| YKL163W | PIR3 | structural constituent of cell wall | -0.4733 | -1.4 | 0.0482333 |
| YBR095C | RXT2 | unknown | -0.4707 | -1.4 | 0.0006189 |
| YOR394W | PAU21 | unknown | -0.4667 | -1.4 | 0.0057981 |
| YAL068C | PAU8 | unknown | -0.4665 | -1.4 | 0.0752435 |
| YIL068C | SEC6 | protein binding | -0.4653 | -1.4 | 0.232778 |
| YGL198W | YIP4 | Rab GTPase binding | -0.4627 | -1.4 | 0.0198333 |
| YNR018W | | unknown | -0.4625 | -1.4 | 0.0884206 |
| YBL113C | | helicase | -0.4620 | -1.4 | 0.0318955 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YCR049C | | | -0.4605 | -1.4 | 0.172441 |
| YGL066W | SGF73 | structural molecule * | -0.4557 | -1.4 | 0.119983 |
| YPL282C | PAU22 | unknown | -0.4543 | -1.4 | 0.0942943 |
| YDR497C | ITR1 | myo-inositol transporter | -0.4523 | -1.4 | 0.0617382 |
| YJL079C | PRY1 | unknown | -0.4522 | -1.4 | 0.0709253 |
| YGL076C | RPL7A | structural constituent of ribosome | -0.4507 | -1.4 | 0.0859879 |
| YFL043C | | | -0.4473 | -1.4 | 0.0095589 |
| YPR165W | RHO1 | GTPase * | -0.4420 | -1.4 | 0.144566 |
| YPR108W | RPN7 | structural molecule | -0.4415 | -1.4 | 0.0871089 |
| YBR301W | DAN3 | unknown | -0.4373 | -1.4 | 0.0778786 |
| YDR077W | SED1 | structural constituent of cell wall | -0.4367 | -1.4 | 0.0248061 |
| YGR295C | COS6 | unknown | -0.4360 | -1.4 | 0.127048 |
| YPR100W | MRPL51 | structural constituent of ribosome | -0.4322 | -1.3 | 0.153402 |
| YJL166W | QCR8 | ubiquinol-cytochrome-c reductase | -0.4295 | -1.3 | 0.181686 |
| YDR080W | VPS41 | Rab guanyl-nucleotide exchange factor | -0.4268 | -1.3 | 0.136035 |
| YOR383C | FIT3 | unknown | -0.4210 | -1.3 | 0.306351 |
| YBR088C | POL30 | DNA polymerase processivity factor | -0.4165 | -1.3 | 0.0010976 |
| YBR088C | POL30 | DNA polymerase processivity factor | -0.4165 | -1.3 | 0.0010976 |
| YPL048W | CAM1 | translation elongation factor | -0.4132 | -1.3 | 0.360113 |
| YCL025C | AGP1 | amino acid transporter | -0.4122 | -1.3 | 0.0445531 |
| YJR023C | | | -0.4122 | -1.3 | 0.0651461 |
| YJL151C | SNA3 | unknown | -0.4105 | -1.3 | 0.0842497 |
| YER067W | | unknown | -0.4103 | -1.3 | 0.0751532 |
| YER103W | SSA4 | ATPase * | -0.4088 | -1.3 | 0.21908 |
| YDR253C | MET32 | DNA binding* | -0.4057 | -1.3 | 0.131526 |
| YER057C | HMF1 | unknown | -0.4050 | -1.3 | 0.0253396 |
| YCR087W | | | -0.4033 | -1.3 | 0.163126 |
| YOR332W | VMA4 | hydrogen-transporting ATPase , rotational mechanism | -0.4032 | -1.3 | 0.161853 |
| YFL067W | | unknown | -0.4018 | -1.3 | 0.596477 |
| YJL073W | JEM1 | unfolded protein binding* | -0.3995 | -1.3 | 0.0082648 |
| YDR251W | PAM1 | unknown | -0.3967 | -1.3 | 0.0264071 |
| YAL003W | EFB1 | translation elongation factor | -0.3927 | -1.3 | 0.0584031 |
| YHR193C | EGD2 | unfolded protein binding | -0.3907 | -1.3 | 0.366134 |
| YPR026W | ATH1 | alpha, alpha-trehalase | -0.3903 | -1.3 | 0.0047912 |
| YIR014W | | unknown | -0.3900 | -1.3 | 0.164385 |
| YMR008C | PLB1 | lysophospholipase | -0.3897 | -1.3 | 0.0161228 |
| YEL011W | GLC3 | 1,4-alpha-glucan branching enzyme | -0.3878 | -1.3 | 0.0152677 |
| YGL112C | TAF6 | general RNA polymerase II transcription factor | -0.3872 | -1.3 | 0.117595 |
| YDL171C | GLT1 | glutamate synthase (NADH) | -0.3857 | -1.3 | 0.263044 |
| YDR329C | PEX3 | protein binding | -0.3857 | -1.3 | 0.0415069 |
| YIR003W | | unknown | -0.3855 | -1.3 | 0.0160999 |
| YKR096W | | unknown | -0.3850 | -1.3 | 0.069603 |
| YJR085C | | unknown | -0.3823 | -1.3 | 0.0481058 |
| YLL024C | SSA2 | ATPase * | -0.3762 | -1.3 | 0.0317893 |
| YIL094C | LYS12 | homoisocitrate dehydrogenase | -0.3730 | -1.3 | 0.0869456 |
| YDL248W | COS7 | receptor | -0.3707 | -1.3 | 0.0216969 |
| YGR142W | BTN2 | unknown | -0.3695 | -1.3 | 0.485944 |
| YLR190W | MMR1 | unknown | -0.3695 | -1.3 | 0.0571431 |
| YHR200W | RPN10 | endopeptidase | -0.3687 | -1.3 | 0.138534 |
| YPL224C | MMT2 | unknown | -0.3682 | -1.3 | 0.110035 |
| YJL223C | PAU1 | unknown | -0.3615 | -1.3 | 0.0897186 |
| YER052C | HOM3 | aspartate kinase | -0.3575 | -1.3 | 0.224273 |
| YDR542W | PAU10 | unknown | -0.3568 | -1.3 | 0.193424 |
| YJL216C | | alpha-galactosidase * | -0.3530 | -1.3 | 0.0213934 |
| YJL216C | | alpha-galactosidase * | -0.3530 | -1.3 | 0.0213934 |
| YKL058W | TOA2 | general RNA polymerase II transcription factor | -0.3465 | -1.3 | 0.0209133 |
| YGR038W | ORM1 | unknown | -0.3463 | -1.3 | 0.0206693 |
| YCR102C | | unknown | -0.3437 | -1.3 | 0.445094 |
| YDL014W | NOP1 | methyltransferase | -0.3407 | -1.3 | 0.107289 |
| YKL013C | ARC19 | structural molecule | -0.3405 | -1.3 | 0.357407 |
| YIL097W | FYV10 | unknown | -0.3335 | -1.3 | 0.199443 |
| YNL104C | LEU4 | 2-isopropylmalate synthase | -0.3332 | -1.3 | 0.353002 |
| YBR036C | CSG2 | enzyme regulator | -0.3270 | -1.3 | 0.0770719 |
| YBR121C | GRS1 | glycine-tRNA ligase | -0.3265 | -1.3 | 0.183405 |
| YDR167W | TAF10 | general RNA polymerase II transcription factor | -0.3260 | -1.3 | 0.174245 |
| YAL061W | | oxidoreductase , acting on the CH-OH group of donors, NAD or NADP | -0.3253 | -1.3 | 0.0522863 |
| YGR135W | PRE9 | endopeptidase | -0.3250 | -1.3 | 0.0800802 |
| YAL062W | GDH3 | glutamate dehydrogenase | -0.3237 | -1.3 | 0.0251359 |
| YPL118W | MRP51 | structural constituent of ribosome | -0.3233 | -1.3 | 0.0106163 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YCR104W | PAU3 | unknown | -0.3228 | -1.3 | 0.0714356 |
| YBR234C | ARC40 | structural constituent of cytoskeleton | -0.3222 | -1.3 | 0.151189 |
| YJL202C | | | -0.3222 | -1.3 | 0.46558 |
| YBR047W | FMP23 | unknown | -0.3205 | -1.2 | 0.0364289 |
| YIL085C | KTR7 | mannosyltransferase | -0.3197 | -1.2 | 0.875058 |
| YFR017C | | unknown | -0.3195 | -1.2 | 0.0218165 |
| YJL072C | PSF2 | unknown | -0.3182 | -1.2 | 0.0391689 |
| YBL066C | SEF1 | unknown | -0.3112 | -1.2 | 0.253791 |
| YOR374W | ALD4 | aldehyde dehydrogenase | -0.3110 | -1.2 | 0.0074353 |
| YCR014C | POL4 | beta DNA polymerase | -0.3105 | -1.2 | 0.193762 |
| YBR202W | CDC47 | chromatin binding* | -0.3097 | -1.2 | 0.0939306 |
| YCL064C | CHA1 | L-serine ammonia-lyase * | -0.3072 | -1.2 | 0.0354441 |
| YJL206C-A | | | -0.3067 | -1.2 | 0.18422 |
| YDR099W | BMH2 | protein binding* | -0.3000 | -1.2 | 0.403096 |
| YDR266C | | unknown | -0.2998 | -1.2 | 0.0253857 |
| YCL051W | LRE1 | transcription regulator * | -0.2995 | -1.2 | 0.342779 |
| YDL213C | NOP6 | unknown | -0.2987 | -1.2 | 0.284312 |
| YKL109W | HAP4 | transcriptional activator | -0.2958 | -1.2 | 0.184835 |
| YBR102C | EXO84 | protein binding | -0.2937 | -1.2 | 0.261787 |
| TDR261C-D | | | -0.2923 | -1.2 | 0.195501 |
| YKL093W | MBR1 | unknown | -0.2897 | -1.2 | 0.0637364 |
| YBR247C | ENP1 | snoRNA binding | -0.2890 | -1.2 | 0.109716 |
| YCL028W | RNQ1 | unknown | -0.2883 | -1.2 | 0.147312 |
| YEL049W | PAU2 | unknown | -0.2850 | -1.2 | 0.145279 |
| YBR052C | RFS1 | unknown | -0.2835 | -1.2 | 0.165765 |
| YMR311C | GLC8 | enzyme activator | -0.2825 | -1.2 | 0.150182 |
| YBR053C | | unknown | -0.2793 | -1.2 | 0.192995 |
| YHL021C | FMP12 | unknown | -0.2720 | -1.2 | 0.421484 |
| YBR218C | PYC2 | pyruvate carboxylase | -0.2677 | -1.2 | 0.198903 |
| YCL001W | RER1 | unknown | -0.2667 | -1.2 | 0.126119 |
| YMR325W | PAU19 | unknown | -0.2662 | -1.2 | 0.580808 |
| YDR156W | RPA14 | DNA-directed RNA polymerase | -0.2658 | -1.2 | 0.246353 |
| YBR287W | ZSP1 | unknown | -0.2625 | -1.2 | 0.817135 |
| YDR496C | PUF6 | specific transcriptional repressor | -0.2617 | -1.2 | 0.261561 |
| YHR026W | PPA1 | hydrogen-transporting ATPase , rotational mechanism | -0.2593 | -1.2 | 0.171852 |
| YKL217W | JEN1 | lactate transporter | -0.2590 | -1.2 | 0.476441 |
| YMR186W | HSC82 | unfolded protein binding | -0.2563 | -1.2 | 0.0485359 |
| YBR263W | SHM1 | glycine hydroxymethyltransferase | -0.2562 | -1.2 | 0.0701914 |
| YBR082C | UBC4 | ubiquitin conjugating enzyme | -0.2553 | -1.2 | 0.158767 |
| YIL066C | RNR3 | ribonucleoside-diphosphate reductase | -0.2550 | -1.2 | 0.142169 |
| YBR015C | MNN2 | alpha-1,2-mannosyltransferase | -0.2532 | -1.2 | 0.39925 |
| YIL084C | SDS3 | protein binding | -0.2532 | -1.2 | 0.260757 |
| YPR028W | YOP1 | protein binding | -0.2532 | -1.2 | 0.333069 |
| YIL113W | SDP1 | MAP kinase | -0.2518 | -1.2 | 0.529468 |
| YBR010W | HHT1 | DNA binding | -0.2512 | -1.2 | 0.14227 |
| YML039W | | | -0.2512 | -1.2 | 0.399605 |
| YMR080C | NAM7 | ATP-dependent RNA helicase * | -0.2507 | -1.2 | 0.362955 |
| YNL022C | | unknown | -0.2500 | -1.2 | 3.492E-05 |
| YJL222W | VTH2 | signal sequence binding | -0.2487 | -1.2 | 0.113229 |
| YBR170C | NPL4 | unknown | -0.2483 | -1.2 | 0.312997 |
| YML012W | ERV25 | unknown | -0.2480 | -1.2 | 0.929802 |
| YHR194W | MDM31 | unknown | -0.2455 | -1.2 | 0.254302 |
| YGR008C | STF2 | unknown | -0.2452 | -1.2 | 0.143857 |
| YLR437C | | unknown | -0.2395 | -1.2 | 0.0535503 |
| YNL209W | SSB2 | ATPase * | -0.2388 | -1.2 | 0.194611 |
| YCL037C | SRO9 | RNA binding | -0.2387 | -1.2 | 0.447294 |
| YIL041W | GVP36 | unknown | -0.2382 | -1.2 | 0.0043406 |
| YKR046C | PET10 | unknown | -0.2365 | -1.2 | 0.322573 |
| YAL067C | SEO1 | transporter | -0.2340 | -1.2 | 0.0472514 |
| YCL008C | STP22 | protein binding | -0.2327 | -1.2 | 0.149481 |
| YDR232W | HEM1 | 5-aminolevulinate synthase | -0.2327 | -1.2 | 0.502076 |
| YHR211W | FLO5 | mannose binding | -0.2325 | -1.2 | 0.354771 |
| YBR149W | ARA1 | aldo-keto reductase * | -0.2317 | -1.2 | 0.157737 |
| YNL336W | COS1 | unknown | -0.2307 | -1.2 | 0.287355 |
| YGR204W | ADE3 | formate-tetrahydrofolate ligase | -0.2293 | -1.2 | 0.363124 |
| YHR183W | GND1 | phosphogluconate dehydrogenase | -0.2247 | -1.2 | 0.025193 |
| YHR213W | | unknown | -0.2192 | -1.2 | 0.112611 |
| YBR180W | DTR1 | multidrug transporter | -0.2173 | -1.2 | 0.793773 |
| YOR141C | ARP8 | unknown | -0.2163 | -1.2 | 0.0915507 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YGR037C | ACB1 | long-chain fatty acid transporter | -0.2142 | -1.2 | 0.199754 |
| YPL190C | NAB3 | poly(A) binding | -0.2138 | -1.2 | 0.324583 |
| YGR248W | SOL4 | 6-phosphogluconolactonase | -0.2112 | -1.2 | 0.174259 |
| YNL160W | YGP1 | unknown | -0.2105 | -1.2 | 0.313319 |
| YDL086W | | carboxymethylenebutenolidase | -0.2093 | -1.2 | 0.253101 |
| YKR094C | RPL40B | structural constituent of ribosome* | -0.2093 | -1.2 | 0.393069 |
| YDR201W | SPC19 | structural constituent of cytoskeleton | -0.2073 | -1.2 | 0.18729 |
| YGR250C | | RNA binding | -0.2070 | -1.2 | 0.229686 |
| YDL184C | RPL41A | structural constituent of ribosome | -0.2065 | -1.2 | 0.17499 |
| YLR055C | SPT8 | transcription cofactor | -0.2043 | -1.2 | 0.0635119 |
| YBR280C | | unknown | -0.2040 | -1.2 | 0.0055926 |
| YLR216C | CPR6 | unfolded protein binding* | -0.2037 | -1.2 | 0.0513457 |
| YIL142W | CCT2 | unfolded protein binding | -0.1995 | -1.1 | 0.0212486 |
| YER080W | FMP29 | unknown | -0.1988 | -1.1 | 0.106618 |
| YHR157W | REC104 | unknown | -0.1977 | -1.1 | 0.254376 |
| YMR196W | | unknown | -0.1958 | -1.1 | 0.469018 |
| YIL156W | UBP7 | ubiquitin-specific protease | -0.1892 | -1.1 | 0.0608192 |
| YBR269C | FMP21 | unknown | -0.1882 | -1.1 | 0.279132 |
| YCL017C | NFS1 | cystathionine gamma-lyase * | -0.1870 | -1.1 | 0.269647 |
| YBR100W | | | -0.1850 | -1.1 | 0.0206199 |
| YBR100W | | | -0.1850 | -1.1 | 0.0206199 |
| YDR310C | SUM1 | transcriptional repressor | -0.1800 | -1.1 | 0.959801 |
| YJL160C | | unknown | -0.1782 | -1.1 | 0.0723843 |
| YBR125C | PTC4 | protein phosphatase type 2C | -0.1777 | -1.1 | 0.63724 |
| YDR427W | RPN9 | structural molecule | -0.1762 | -1.1 | 0.517973 |
| YPL085W | SEC16 | structural molecule | -0.1757 | -1.1 | 0.553842 |
| YPL105C | | unknown | -0.1722 | -1.1 | 0.438361 |
| YKL171W | | protein kinase | -0.1705 | -1.1 | 0.47087 |
| YFR031C-A | RPL2A | structural constituent of ribosome | -0.1687 | -1.1 | 0.0372032 |
| YOR375C | GDH1 | glutamate dehydrogenase | -0.1675 | -1.1 | 0.11877 |
| YCL019W | | | -0.1650 | -1.1 | 0.0460387 |
| YIL050W | PCL7 | cyclin-dependent protein kinase | -0.1650 | -1.1 | 0.444316 |
| YPR033C | HTS1 | histidine-tRNA ligase | -0.1650 | -1.1 | 0.0420972 |
| YOR185C | GSP2 | GTPase | -0.1632 | -1.1 | 0.215822 |
| YCR082W | AHC2 | unknown | -0.1618 | -1.1 | 0.103037 |
| YOR372C | NDD1 | transcriptional activator | -0.1615 | -1.1 | 0.0196284 |
| YJL210W | PEX2 | protein binding | -0.1607 | -1.1 | 0.378566 |
| YPL240C | HSP82 | ATPase , coupled | -0.1603 | -1.1 | 0.0459967 |
| YBR009C | HHF1 | DNA binding | -0.1588 | -1.1 | 0.341646 |
| YBR069C | TAT1 | amino acid transporter | -0.1557 | -1.1 | 0.599037 |
| YNL134C | | alcohol dehydrogenase | -0.1557 | -1.1 | 0.762409 |
| YDR088C | SLU7 | RNA splicing factor | -0.1552 | -1.1 | 0.143022 |
| YDR180W | SCC2 | unknown | -0.1528 | -1.1 | 0.0995337 |
| YIL075C | RPN2 | endopeptidase * | -0.1495 | -1.1 | 0.249126 |
| YJL101C | GSH1 | glutamate-cysteine ligase | -0.1427 | -1.1 | 0.0886188 |
| YCL027W | FUS1 | unknown | -0.1405 | -1.1 | 0.155886 |
| YIL162W | SUC2 | beta-fructofuranosidase | -0.1400 | -1.1 | 0.544858 |
| YHR186C | KOG1 | unknown | -0.1350 | -1.1 | 0.764435 |
| YJL123C | | unknown | -0.1345 | -1.1 | 0.965833 |
| YBR150C | TBS1 | unknown | -0.1332 | -1.1 | 0.315707 |
| YLR256W | HAP1 | specific RNA polymerase II transcription factor | -0.1332 | -1.1 | 0.544145 |
| YAL030W | SNC1 | v-SNARE | -0.1295 | -1.1 | 0.718942 |
| YAL011W | SWC3 | unknown | -0.1288 | -1.1 | 0.16215 |
| YAL035W | FUN12 | GTPase * | -0.1283 | -1.1 | 0.0259497 |
| YJL020C | BBC1 | myosin I binding | -0.1268 | -1.1 | 0.552426 |
| YLR158C | ASP3-3 | asparaginase | -0.1257 | -1.1 | 0.245713 |
| YOR123C | LEO1 | RNA polymerase II transcription elongation factor | -0.1247 | -1.1 | 0.857463 |
| YJR015W | | unknown | -0.1240 | -1.1 | 0.742664 |
| YKR048C | NAP1 | protein binding | -0.1240 | -1.1 | 0.146526 |
| YCR021C | HSP30 | unknown | -0.1225 | -1.1 | 0.445803 |
| YCRX21C | | | -0.1225 | -1.1 | 0.196101 |
| YJR105W | ADO1 | adenosine kinase | -0.1222 | -1.1 | 0.887085 |
| YGL148W | ARO2 | chorismate synthase * | -0.1215 | -1.1 | 0.580228 |
| YFR033C | QCR6 | ubiquinol-cytochrome-c reductase | -0.1203 | -1.1 | 0.352352 |
| YBR289W | SNF5 | general RNA polymerase II transcription factor | -0.1192 | -1.1 | 0.71639 |
| YFL066C | | unknown | -0.1167 | -1.1 | 0.983209 |
| YJL021C | | | -0.1165 | -1.1 | 0.19242 |
| YAL042W | ERV46 | unknown | -0.1153 | -1.1 | 0.0072213 |
| YNL030W | HHF2 | DNA binding | -0.1147 | -1.1 | 0.213936 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YEL047C | | fumarate reductase (NADH) | -0.1133 | -1.1 | 0.038208 |
| YHR217C | | | -0.1130 | -1.1 | 0.245118 |
| YIL154C | IMP2' | transcription coactivator | -0.1125 | -1.1 | 0.623778 |
| YBR255W | | unknown | -0.1123 | -1.1 | 0.673023 |
| YKL150W | MCR1 | cytochrome-b5 reductase | -0.1112 | -1.1 | 0.191252 |
| YBR159W | IFA38 | ketoreductase | -0.1103 | -1.1 | 0.432728 |
| YCRX13W | | | -0.1093 | -1.1 | 0.187926 |
| YAL029C | MYO4 | microfilament motor | -0.1092 | -1.1 | 0.716549 |
| YHR158C | KEL1 | unknown | -0.1073 | -1.1 | 0.638851 |
| YCR046C | IMG1 | structural constituent of ribosome | -0.1068 | -1.1 | 0.872004 |
| YPR002W | PDH1 | unknown | -0.1068 | -1.1 | 0.640189 |
| YHR179W | OYE2 | NADPH dehydrogenase | -0.1057 | -1.1 | 0.521314 |
| YLR204W | QRI5 | unknown | -0.1052 | -1.1 | 0.798823 |
| YGR256W | GND2 | phosphogluconate dehydrogenase | -0.1048 | -1.1 | 0.700165 |
| YLR247C | | helicase | -0.1032 | -1.1 | 0.156079 |
| YBR008C | FLR1 | multidrug transporter | -0.1005 | -1.1 | 0.133196 |
| YAL016W | TPD3 | protein phosphatase type 2A | -0.0998 | -1.1 | 0.779859 |
| YOL039W | RPP2A | structural constituent of ribosome | -0.0957 | -1.1 | 0.440778 |
| YIL005W | EPS1 | protein disulfide isomerase | -0.0917 | -1.1 | 0.688447 |
| YBR156C | SLI15 | protein kinase | -0.0892 | -1.1 | 0.639801 |
| YBR143C | SUP45 | translation release factor | -0.0885 | -1.1 | 0.202823 |
| YBR205W | KTR3 | mannosyltransferase | -0.0882 | -1.1 | 0.0554639 |
| YJL130C | URA2 | carbamoyl-phosphate synthase (glutamine-hydrolyzing) * | -0.0873 | -1.1 | 0.830579 |
| YBR245C | ISW1 | ATPase | -0.0872 | -1.1 | 0.621223 |
| YDR059C | UBC5 | ubiquitin conjugating enzyme | -0.0848 | -1.1 | 0.837788 |
| YER043C | SAH1 | adenosylhomocysteinase | -0.0847 | -1.1 | 0.0318519 |
| YMR020W | FMS1 | amine oxidase | -0.0835 | -1.1 | 0.767023 |
| YJL070C | | unknown | -0.0822 | -1.1 | 0.37809 |
| YNL031C | HHT2 | DNA binding | -0.0813 | -1.1 | 0.807911 |
| YDR154C | | | -0.0808 | -1.1 | 0.346086 |
| YKL056C | TMA19 | unknown | -0.0778 | -1.1 | 0.569459 |
| YBR126C | TPS1 | alpha, alpha-trehalose-phosphate synthase (UDP-forming) | -0.0747 | -1.1 | 0.090745 |
| YOR327C | SNC2 | v-SNARE | -0.0732 | -1.1 | 0.79496 |
| YDL140C | RPO21 | DNA-directed RNA polymerase | -0.0720 | -1.1 | 0.620539 |
| YBR239C | | unknown | -0.0710 | -1.1 | 0.222801 |
| YMR226C | TMA29 | oxidoreductase, acting on the aldehyde or oxo group of donors, N | -0.0660 | -1.0 | 0.390916 |
| YPL234C | TFP3 | hydrogen-transporting ATPase, rotational mechanism | -0.0632 | -1.0 | 0.998273 |
| YKR002W | PAP1 | polynucleotide adenyltransferase | -0.0617 | -1.0 | 0.78612 |
| YAR042W | SWH1 | phosphatidylinositol binding | -0.0597 | -1.0 | 0.881636 |
| YBR109C | CMD1 | protein binding* | -0.0592 | -1.0 | 0.422345 |
| YGL221C | NIF3 | unknown | -0.0588 | -1.0 | 0.952809 |
| YDL220C | CDC13 | single-stranded DNA binding | -0.0587 | -1.0 | 0.888505 |
| YML100W | TSL1 | enzyme regulator * | -0.0580 | -1.0 | 0.994651 |
| YLR299W | ECM38 | protein-glutamine gamma-glutamyltransferase | -0.0572 | -1.0 | 0.768608 |
| YOR291W | | unknown | -0.0560 | -1.0 | 0.802987 |
| YBR071W | | unknown | -0.0558 | -1.0 | 0.901496 |
| YML010W | SPT5 | RNA polymerase II transcription elongation factor | -0.0553 | -1.0 | 0.671085 |
| YLR436C | ECM30 | unknown | -0.0535 | -1.0 | 0.623946 |
| YJL221C | FSP2 | alpha-glucosidase | -0.0518 | -1.0 | 0.656974 |
| YEL053C | MAK10 | peptide alpha-N-acetyltransferase | -0.0510 | -1.0 | 0.867231 |
| YOR120W | GCY1 | aldo-keto reductase | -0.0508 | -1.0 | 0.56991 |
| YJL142C | | | -0.0502 | -1.0 | 0.336986 |
| YMR212C | EFR3 | unknown | -0.0467 | -1.0 | 0.976364 |
| YBR137W | | unknown | -0.0465 | -1.0 | 0.824171 |
| YJR007W | SUI2 | translation initiation factor | -0.0430 | -1.0 | 0.256203 |
| YKR071C | DRE2 | unknown | -0.0408 | -1.0 | 0.555714 |
| YBR079C | RPG1 | translation initiation factor | -0.0375 | -1.0 | 0.91121 |
| YJL149W | | unknown | -0.0372 | -1.0 | 0.925782 |
| YLR350W | ORM2 | unknown | -0.0368 | -1.0 | 0.758815 |
| YER068W | MOT2 | 3'-5'-exoribonuclease * | -0.0365 | -1.0 | 0.678687 |
| YCL034W | LSB5 | unknown | -0.0358 | -1.0 | 0.439616 |
| YKL142W | MRP8 | structural constituent of ribosome | -0.0358 | -1.0 | 0.342182 |
| YBR132C | AGP2 | amino acid transporter | -0.0352 | -1.0 | 0.778329 |
| YNL305C | | unknown | -0.0343 | -1.0 | 0.635317 |
| YJL115W | ASF1 | transcription regulator * | -0.0332 | -1.0 | 0.467758 |
| YDR513W | GRX2 | thiol-disulfide exchange intermediate * | -0.0328 | -1.0 | 0.74735 |
| YLR249W | YEF3 | translation elongation factor | -0.0323 | -1.0 | 0.538453 |
| YPR149W | NCE102 | unknown | -0.0317 | -1.0 | 0.409441 |
| YDR091C | RLI1 | ATPase * | -0.0305 | -1.0 | 0.993499 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YJL016W | | unknown | -0.0302 | -1.0 | 0.631493 |
| YER026C | CHO1 | CDP-diacylglycerol-serine O-phosphatidyltransferase | -0.0283 | -1.0 | 0.966644 |
| YLR270W | DCS1 | hydrolase, acting on acid anhydrides, in phosphorus-containing a | -0.0278 | -1.0 | 0.793073 |
| YJR104C | SOD1 | copper, zinc superoxide dismutase | -0.0265 | -1.0 | 0.82633 |
| YAR014C | BUD14 | protein phosphatase type 1 regulator | -0.0257 | -1.0 | 0.477668 |
| YMR105C | PGM2 | phosphoglucosmutase | -0.0252 | -1.0 | 0.870963 |
| YDR250C | | | -0.0243 | -1.0 | 0.092452 |
| YER072W | VTC1 | unknown | -0.0237 | -1.0 | 0.626125 |
| YBL041W | PRE7 | endopeptidase | -0.0205 | -1.0 | 0.566421 |
| YHR131C | | unknown | -0.0200 | -1.0 | 0.536315 |
| YPR080W | TEF1 | translation elongation factor | -0.0187 | -1.0 | 0.769361 |
| YDL005C | MED2 | RNA polymerase II transcription mediator | -0.0137 | -1.0 | 0.595136 |
| YFR016C | | unknown | -0.0123 | -1.0 | 0.695092 |
| YOL129W | VPS68 | unknown | -0.0110 | -1.0 | 0.104815 |
| YOR285W | | unknown | -0.0088 | -1.0 | 0.945477 |
| YOR108W | LEU9 | 2-isopropylmalate synthase | -0.0082 | -1.0 | 0.939298 |
| YNL016W | PUB1 | mRNA binding* | -0.0078 | -1.0 | 0.924079 |
| YBR177C | EHT1 | serine hydrolase | -0.0075 | -1.0 | 0.93439 |
| YDR483W | KRE2 | alpha-1,2-mannosyltransferase | -0.0073 | -1.0 | 0.191345 |
| YLR157C | ASP3-2 | asparaginase | -0.0073 | -1.0 | 0.869328 |
| YHR096C | HXT5 | glucose transporter | -0.0048 | -1.0 | 0.671767 |
| YCR042C | TAF2 | general RNA polymerase II transcription factor | -0.0043 | -1.0 | 0.774726 |
| YJL140W | RPB4 | DNA-directed RNA polymerase | -0.0043 | -1.0 | 0.323239 |
| YJL095W | BCK1 | MAP kinase | -0.0042 | -1.0 | 0.792483 |
| YHR190W | ERG9 | farnesyl-diphosphate farnesyltransferase | -0.0038 | -1.0 | 0.71858 |
| YBR090C | | unknown | -0.0030 | -1.0 | 0.547006 |
| YBL097W | BRN1 | unknown | -0.0027 | -1.0 | 0.649356 |
| YBR206W | | | -0.0020 | -1.0 | 0.0744164 |
| YDR520C | | unknown | -0.0007 | -1.0 | 0.826762 |
| YGL243W | TAD1 | tRNA specific adenosine deaminase | -0.0007 | -1.0 | 0.31548 |
| YJL156C | SSY5 | amino acid binding | 0.0030 | 1.0 | 0.773274 |
| YBR115C | LYS2 | L-aminoadipate-semialdehyde dehydrogenase | 0.0032 | 1.0 | 0.469014 |
| YDL099W | BUG1 | unknown | 0.0040 | 1.0 | 0.330018 |
| YPR062W | FCY1 | cytosine deaminase | 0.0068 | 1.0 | 0.808286 |
| YPL274W | SAM3 | S-adenosylmethionine transporter | 0.0070 | 1.0 | 0.866622 |
| YDL095W | PMT1 | dolichyl-phosphate-mannose-protein mannosyltransferase | 0.0077 | 1.0 | 0.868196 |
| YDL174C | DLD1 | D-lactate dehydrogenase | 0.0087 | 1.0 | 0.0028314 |
| YPL222W | FMP40 | unknown | 0.0090 | 1.0 | 0.873304 |
| YLR246W | ERF2 | protein-cysteine S-palmitoleyltransferase | 0.0093 | 1.0 | 0.0984154 |
| YDR054C | CDC34 | ubiquitin conjugating enzyme | 0.0095 | 1.0 | 0.334444 |
| YOR053W | | | 0.0097 | 1.0 | 0.491963 |
| YJL092W | HPR5 | DNA helicase | 0.0113 | 1.0 | 0.34034 |
| YLR160C | ASP3-4 | asparaginase | 0.0127 | 1.0 | 0.947893 |
| YML074C | FPR3 | peptidyl-prolyl cis-trans isomerase | 0.0148 | 1.0 | 0.405274 |
| YDR316W-B | | | 0.0150 | 1.0 | 0.862321 |
| YAL039C | CYC3 | holocytochrome-c synthase | 0.0157 | 1.0 | 0.657798 |
| YBR262C | FMP51 | unknown | 0.0173 | 1.0 | 0.881224 |
| YDL131W | LYS21 | homocitrate synthase | 0.0180 | 1.0 | 0.902757 |
| YBR118W | TEF2 | translation elongation factor | 0.0188 | 1.0 | 0.68477 |
| YBR236C | ABD1 | mRNA (guanine-N7-)-methyltransferase | 0.0208 | 1.0 | 0.942208 |
| YBL081W | | unknown | 0.0223 | 1.0 | 0.369428 |
| YLR155C | ASP3-1 | asparaginase | 0.0232 | 1.0 | 0.0389851 |
| YBR240C | THI2 | transcriptional activator | 0.0237 | 1.0 | 0.818587 |
| YPL265W | DIP5 | amino acid transporter | 0.0243 | 1.0 | 0.354954 |
| YHR009C | | unknown | 0.0268 | 1.0 | 0.383037 |
| YCR044C | PER1 | unknown | 0.0277 | 1.0 | 0.745013 |
| YLR381W | CTF3 | protein binding | 0.0305 | 1.0 | 0.392949 |
| YMR302C | YME2 | exonuclease | 0.0323 | 1.0 | 0.0942745 |
| YCRX17W | | | 0.0360 | 1.0 | 0.883746 |
| YBR251W | MRPS5 | structural constituent of ribosome | 0.0368 | 1.0 | 0.733327 |
| YJL225C | | helicase | 0.0387 | 1.0 | 0.412489 |
| YFL016C | MDJ1 | unfolded protein binding* | 0.0388 | 1.0 | 0.812296 |
| YBR237W | PRP5 | RNA splicing factor | 0.0393 | 1.0 | 0.788099 |
| YCR053W | THR4 | threonine synthase | 0.0413 | 1.0 | 0.841583 |
| YDR473C | PRP3 | RNA splicing factor | 0.0430 | 1.0 | 0.985185 |
| YBR070C | ALG14 | N-acetylglucosaminyldiphosphodolichol N-acetylglucosaminyltra | 0.0438 | 1.0 | 0.90828 |
| YFL008W | SMC1 | ATPase * | 0.0465 | 1.0 | 0.586551 |
| YCL041C | | | 0.0488 | 1.0 | 0.797544 |
| YDL124W | | alpha-keto amide reductase * | 0.0505 | 1.0 | 0.971075 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YDL161W | ENT1 | clathrin binding | 0.0505 | 1.0 | 0.759655 |
| YJR002W | MPP10 | unknown | 0.0507 | 1.0 | 0.353355 |
| YOR304W | ISW2 | ATPase | 0.0507 | 1.0 | 0.806684 |
| YBR250W | | unknown | 0.0523 | 1.0 | 0.453664 |
| YBR119W | MUD1 | RNA binding | 0.0525 | 1.0 | 0.770962 |
| YGL227W | VID30 | unknown | 0.0533 | 1.0 | 0.897366 |
| YDR208W | MSS4 | 1-phosphatidylinositol-4-phosphate 5-kinase | 0.0545 | 1.0 | 0.0877582 |
| YKR042W | UTH1 | unknown | 0.0547 | 1.0 | 0.7362 |
| YBR011C | IPP1 | inorganic diphosphatase | 0.0553 | 1.0 | 0.542188 |
| YCR032W | BPH1 | unknown | 0.0567 | 1.0 | 0.978016 |
| YBR208C | DUR1,2 | allophanate hydrolase * | 0.0568 | 1.0 | 0.786421 |
| YCL003W | | | 0.0573 | 1.0 | 0.675468 |
| YLR178C | TFS1 | lipid binding* | 0.0573 | 1.0 | 0.924812 |
| YEL027W | CUP5 | hydrogen ion transporter | 0.0600 | 1.0 | 0.935443 |
| YFR008W | FAR7 | unknown | 0.0600 | 1.0 | 0.0528463 |
| YJL174W | KRE9 | unknown | 0.0608 | 1.0 | 0.625319 |
| YDL186W | | unknown | 0.0615 | 1.0 | 0.819761 |
| YOR113W | AZF1 | DNA binding* | 0.0628 | 1.0 | 0.0594089 |
| YGR155W | CYS4 | cystathionine beta-synthase | 0.0638 | 1.0 | 0.710362 |
| YBR024W | SCO2 | thioredoxin peroxidase | 0.0640 | 1.0 | 0.4598 |
| YNL102W | POL1 | alpha DNA polymerase | 0.0670 | 1.0 | 0.552319 |
| YDR309C | GIC2 | small GTPase regulator | 0.0710 | 1.1 | 0.957103 |
| YOR182C | RPS30B | structural constituent of ribosome | 0.0718 | 1.1 | 0.957406 |
| YJL138C | TIF2 | translation initiation factor | 0.0723 | 1.1 | 0.330517 |
| YIL121W | QDR2 | multidrug efflux pump | 0.0738 | 1.1 | 0.904926 |
| YDR433W | | | 0.0745 | 1.1 | 0.627933 |
| YDR145W | TAF12 | general RNA polymerase II transcription factor | 0.0760 | 1.1 | 0.094814 |
| YKL107W | | unknown | 0.0762 | 1.1 | 0.882871 |
| YER073W | ALD5 | 3-chloroallyl aldehyde dehydrogenase | 0.0773 | 1.1 | 0.113367 |
| YBR212W | NGR1 | RNA binding* | 0.0775 | 1.1 | 0.607473 |
| YFL058W | THI5 | unknown | 0.0797 | 1.1 | 0.589852 |
| YKL212W | SAC1 | inositol or phosphatidylinositol phosphatase | 0.0800 | 1.1 | 0.714299 |
| YLR300W | EXG1 | glucan 1,3-beta-glucosidase | 0.0808 | 1.1 | 0.109759 |
| YJL209W | CBP1 | mRNA binding | 0.0820 | 1.1 | 0.479526 |
| YEL020C | | unknown | 0.0845 | 1.1 | 0.363063 |
| YDL106C | PHO2 | transcription factor | 0.0858 | 1.1 | 0.687536 |
| YPL206C | | glycerophosphodiester phosphodiesterase | 0.0870 | 1.1 | 0.490154 |
| YIL036W | CST6 | specific RNA polymerase II transcription factor | 0.0873 | 1.1 | 0.704021 |
| YDR330W | UBX5 | unknown | 0.0885 | 1.1 | 0.0172307 |
| YKR090W | PXL1 | protein binding | 0.0932 | 1.1 | 0.283644 |
| YER035W | EDC2 | RNA binding | 0.0955 | 1.1 | 0.247111 |
| YKR059W | TIF1 | translation initiation factor | 0.0980 | 1.1 | 0.681532 |
| YML058W | SML1 | enzyme inhibitor | 0.0997 | 1.1 | 0.108866 |
| YDR127W | ARO1 | 3-dehydroquinate dehydratase * | 0.0998 | 1.1 | 0.0940223 |
| YGL237C | HAP2 | transcriptional activator | 0.1000 | 1.1 | 0.316126 |
| YGR189C | CRH1 | unknown | 0.1027 | 1.1 | 0.302257 |
| YBR224W | | | 0.1055 | 1.1 | 0.594227 |
| YJL129C | TRK1 | potassium ion transporter | 0.1063 | 1.1 | 0.645302 |
| YLR264W | RPS28B | structural constituent of ribosome | 0.1073 | 1.1 | 0.673035 |
| YDR200C | VPS64 | unknown | 0.1077 | 1.1 | 0.749164 |
| YIL026C | IRR1 | protein binding | 0.1102 | 1.1 | 0.57524 |
| YPL260W | | unknown | 0.1113 | 1.1 | 0.347296 |
| YNL023C | FAP1 | transcription factor | 0.1117 | 1.1 | 0.828141 |
| YJL164C | TPK1 | protein serine/threonine kinase | 0.1120 | 1.1 | 0.0311745 |
| YGR118W | RPS23A | structural constituent of ribosome | 0.1140 | 1.1 | 0.731682 |
| YER019C-A | SBH2 | protein transporter | 0.1143 | 1.1 | 0.28505 |
| YLR325C | RPL38 | structural constituent of ribosome | 0.1153 | 1.1 | 0.562779 |
| YBR261C | | S-adenosylmethionine-dependent methyltransferase | 0.1172 | 1.1 | 0.609125 |
| YNL161W | CBK1 | protein kinase | 0.1173 | 1.1 | 0.0073165 |
| YCR080W | | | 0.1177 | 1.1 | 0.0556888 |
| YFL059W | SNZ3 | protein binding | 0.1183 | 1.1 | 0.499157 |
| YGR038C-A | | | 0.1188 | 1.1 | 0.297821 |
| YJR127C | RSF2 | transcription factor | 0.1207 | 1.1 | 0.341464 |
| YAL019W | FUN30 | unknown | 0.1227 | 1.1 | 0.752596 |
| YOL109W | ZEO1 | unknown | 0.1242 | 1.1 | 0.174605 |
| YBR275C | RIF1 | protein binding* | 0.1250 | 1.1 | 0.67596 |
| YDL081C | RPP1A | structural constituent of ribosome | 0.1252 | 1.1 | 0.52939 |
| YDR510W | SMT3 | protein tag | 0.1260 | 1.1 | 0.123016 |
| YLR238W | FAR10 | unknown | 0.1260 | 1.1 | 0.166042 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YEL044W | IES6 | unknown | 0.1262 | 1.1 | 0.277265 |
| YER160C | | | 0.1272 | 1.1 | 0.0450691 |
| YML045W | | | 0.1285 | 1.1 | 0.861791 |
| YDR487C | RIB3 | 3,4-dihydroxy-2-butanone-4-phosphate synthase | 0.1288 | 1.1 | 0.588451 |
| YLR410W-B | | | 0.1313 | 1.1 | 0.346615 |
| YBR140C | IRA1 | Ras GTPase activator | 0.1333 | 1.1 | 0.633536 |
| YBR260C | RGD1 | Rho GTPase activator | 0.1343 | 1.1 | 0.203981 |
| YLR206W | ENT2 | clathrin binding | 0.1352 | 1.1 | 0.096404 |
| YHR039BC | | | 0.1375 | 1.1 | 0.433079 |
| YJR066W | TOR1 | protein binding* | 0.1378 | 1.1 | 0.797059 |
| YBR259W | | unknown | 0.1392 | 1.1 | 0.290645 |
| YDR514C | | unknown | 0.1393 | 1.1 | 0.449497 |
| YLR454W | FMP27 | unknown | 0.1403 | 1.1 | 0.428059 |
| YBR081C | SPT7 | structural molecule | 0.1405 | 1.1 | 0.963621 |
| YPL106C | SSE1 | ATP binding* | 0.1432 | 1.1 | 0.761594 |
| YBR169C | SSE2 | unknown | 0.1447 | 1.1 | 0.18804 |
| YJL111W | CCT7 | unfolded protein binding | 0.1478 | 1.1 | 0.294965 |
| YFL021C-A | | | 0.1488 | 1.1 | 0.60639 |
| YOR321W | PMT3 | dolichyl-phosphate-mannose-protein mannosyltransferase | 0.1488 | 1.1 | 0.3494 |
| YIL130W | ASG1 | unknown | 0.1490 | 1.1 | 0.341437 |
| YPL111W | CAR1 | zinc ion binding* | 0.1503 | 1.1 | 0.0667393 |
| YBR135W | CKS1 | protein kinase | 0.1523 | 1.1 | 0.347538 |
| YBR204C | | serine hydrolase | 0.1523 | 1.1 | 0.435179 |
| YPR023C | EAF3 | histone acetyltransferase | 0.1530 | 1.1 | 0.0414963 |
| YAL017W | PSK1 | protein serine/threonine kinase | 0.1537 | 1.1 | 0.0515194 |
| YIL047C | SYG1 | unknown | 0.1545 | 1.1 | 0.532192 |
| YHR178W | STB5 | transcription factor | 0.1547 | 1.1 | 0.6834 |
| YER109C | FLO8 | specific RNA polymerase II transcription factor | 0.1550 | 1.1 | 0.167306 |
| YBR127C | VMA2 | hydrogen-transporting ATPase , rotational mechanism | 0.1568 | 1.1 | 0.0394191 |
| YDL156W | | unknown | 0.1568 | 1.1 | 0.338632 |
| YBR142W | MAK5 | ATP-dependent RNA helicase | 0.1572 | 1.1 | 0.129344 |
| YDL097C | RPN6 | structural molecule | 0.1575 | 1.1 | 0.0220909 |
| YIL105C | SLM1 | phosphoinositide binding | 0.1600 | 1.1 | 0.0047103 |
| YMR168C | CEP3 | DNA bending * | 0.1618 | 1.1 | 0.127399 |
| YLR301W | | unknown | 0.1620 | 1.1 | 0.895445 |
| YMR241W | YHM2 | DNA binding* | 0.1632 | 1.1 | 0.124467 |
| YNL126W | SPC98 | structural constituent of cytoskeleton | 0.1662 | 1.1 | 0.548123 |
| YBR238C | | unknown | 0.1673 | 1.1 | 0.30045 |
| YLR093C | NYV1 | v-SNARE | 0.1675 | 1.1 | 0.910825 |
| YIL090W | ICE2 | unknown | 0.1680 | 1.1 | 0.181399 |
| YIL104C | SHQ1 | unknown | 0.1692 | 1.1 | 0.770929 |
| YOR128C | ADE2 | phosphoribosylaminoimidazole carboxylase | 0.1695 | 1.1 | 0.712344 |
| YDR324C | UTP4 | snoRNA binding | 0.1712 | 1.1 | 0.116058 |
| YMR046C | | | 0.1725 | 1.1 | 0.642339 |
| YOR315W | SFG1 | specific RNA polymerase II transcription factor | 0.1770 | 1.1 | 0.389039 |
| YCR047C | BUD23 | S-adenosylmethionine-dependent methyltransferase | 0.1808 | 1.1 | 0.196023 |
| YGL151W | NUT1 | unknown | 0.1818 | 1.1 | 0.894705 |
| YNL099C | OCA1 | protein tyrosine phosphatase | 0.1842 | 1.1 | 0.315649 |
| YCR025C | | | 0.1855 | 1.1 | 0.517081 |
| YAL054C | ACS1 | acetate-CoA ligase | 0.1862 | 1.1 | 0.262764 |
| YBR157C | ICS2 | unknown | 0.1865 | 1.1 | 0.603056 |
| YER117W | RPL23B | structural constituent of ribosome | 0.1885 | 1.1 | 0.350012 |
| YMR273C | ZDS1 | protein binding | 0.1887 | 1.1 | 0.423295 |
| YBR227C | MCX1 | unfolded protein binding* | 0.1888 | 1.1 | 0.0966638 |
| YHR135C | YCK1 | casein kinase | 0.1888 | 1.1 | 0.0757577 |
| YMR016C | SOK2 | transcription factor | 0.1902 | 1.1 | 0.021014 |
| YAL002W | VPS8 | unknown | 0.1927 | 1.1 | 0.0104272 |
| YAL002W | VPS8 | unknown | 0.1927 | 1.1 | 0.0104272 |
| YLR327C | TMA10 | unknown | 0.1930 | 1.1 | 0.0147312 |
| YFR050C | PRE4 | endopeptidase | 0.1942 | 1.1 | 0.217265 |
| YBL101W-A | | | 0.1950 | 1.1 | 0.652759 |
| YPR132W | RPS23B | structural constituent of ribosome | 0.1950 | 1.1 | 0.230334 |
| YKR066C | CCP1 | cytochrome-c peroxidase | 0.1967 | 1.1 | 0.432107 |
| YJR113C | RSM7 | structural constituent of ribosome | 0.1968 | 1.1 | 0.0219174 |
| YJR113C | RSM7 | structural constituent of ribosome | 0.1968 | 1.1 | 0.0219174 |
| YMR002W | | unknown | 0.1982 | 1.1 | 0.180848 |
| YBR286W | APE3 | aminopeptidase | 0.2000 | 1.1 | 0.0138792 |
| YDR516C | EMI2 | unknown | 0.2002 | 1.1 | 0.215576 |
| YGR014W | MSB2 | osmosensor | 0.2022 | 1.2 | 0.411294 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YBR181C | RPS6B | structural constituent of ribosome | 0.2027 | 1.2 | 0.330238 |
| YOR167C | RPS28A | structural constituent of ribosome | 0.2050 | 1.2 | 0.635452 |
| YBR136W | MEC1 | protein kinase | 0.2052 | 1.2 | 0.601221 |
| YJR059W | PTK2 | protein kinase | 0.2053 | 1.2 | 0.271615 |
| YMR230W | RPS10B | structural constituent of ribosome | 0.2070 | 1.2 | 0.273742 |
| YML040W | | | 0.2088 | 1.2 | 0.156966 |
| YIL107C | PFK26 | 6-phosphofructo-2-kinase | 0.2090 | 1.2 | 0.0389223 |
| YLR227W-A | | | 0.2092 | 1.2 | 0.691964 |
| YDL084W | SUB2 | protein binding* | 0.2107 | 1.2 | 0.258949 |
| YDR179W-A | | unknown | 0.2118 | 1.2 | 0.0062116 |
| YDR179W-A | | unknown | 0.2118 | 1.2 | 0.0062116 |
| YGL126W | SCS3 | unknown | 0.2155 | 1.2 | 0.563394 |
| YJL064W | | | 0.2158 | 1.2 | 0.0972282 |
| YMR250W | GAD1 | glutamate decarboxylase | 0.2198 | 1.2 | 0.399388 |
| YCLX06C | | | 0.2237 | 1.2 | 0.153075 |
| YBR134W | | | 0.2242 | 1.2 | 0.81739 |
| YCL002C | | unknown | 0.2247 | 1.2 | 0.0888468 |
| YIL112W | HOS4 | NAD-dependent histone deacetylase * | 0.2270 | 1.2 | 0.722533 |
| YOR317W | FAA1 | long-chain-fatty-acid-CoA ligase | 0.2278 | 1.2 | 0.0493882 |
| YJL176C | SWI3 | general RNA polymerase II transcription factor | 0.2287 | 1.2 | 0.141799 |
| YLR027C | AAT2 | aspartate transaminase | 0.2317 | 1.2 | 0.0294767 |
| YBR207W | FTH1 | iron ion transporter | 0.2327 | 1.2 | 0.0993296 |
| YBR120C | CBP6 | unknown | 0.2333 | 1.2 | 0.0739784 |
| YDR244W | PEX5 | peroxisome targeting sequence binding* | 0.2360 | 1.2 | 0.149682 |
| YAR003W | SWD1 | histone lysine N-methyltransferase (H3-K4 specific) | 0.2365 | 1.2 | 0.15794 |
| YAR010C | | | 0.2373 | 1.2 | 0.479165 |
| YAR028W | | unknown | 0.2397 | 1.2 | 0.254112 |
| YNR002C | ATO2 | ammonium transporter | 0.2435 | 1.2 | 0.316446 |
| YCR036W | RBK1 | ATP binding* | 0.2453 | 1.2 | 0.174654 |
| YLL010C | PSR1 | phosphoprotein phosphatase | 0.2460 | 1.2 | 0.149773 |
| YJL204C | RCY1 | protein binding | 0.2462 | 1.2 | 0.414138 |
| YEL012W | UBC8 | ubiquitin conjugating enzyme | 0.2495 | 1.2 | 0.202459 |
| YBR230C | OM14 | unknown | 0.2497 | 1.2 | 0.111798 |
| YMR096W | SNZ1 | protein binding | 0.2518 | 1.2 | 0.5897 |
| YLR048W | RPS0B | structural constituent of ribosome | 0.2535 | 1.2 | 0.109574 |
| YOL011W | PLB3 | lysophospholipase | 0.2535 | 1.2 | 0.287428 |
| YDL125C | HNT1 | hydrolase * | 0.2558 | 1.2 | 0.135089 |
| YLR109W | AHP1 | thioredoxin peroxidase | 0.2577 | 1.2 | 0.253213 |
| YMR110C | HFD1 | 3-chloroalyl aldehyde dehydrogenase | 0.2580 | 1.2 | 0.444213 |
| YGR027C | RPS25A | structural constituent of ribosome | 0.2583 | 1.2 | 0.219048 |
| YDR083W | RRP8 | methyltransferase | 0.2587 | 1.2 | 0.0396527 |
| YDR083W | RRP8 | methyltransferase | 0.2587 | 1.2 | 0.0396527 |
| YLR104W | | unknown | 0.2593 | 1.2 | 0.0771835 |
| YIL045W | PIG2 | protein phosphatase type 1 regulator | 0.2612 | 1.2 | 0.377468 |
| YOL106W | | | 0.2628 | 1.2 | 0.168994 |
| YIR016W | | unknown | 0.2632 | 1.2 | 0.450964 |
| YAL014C | SYN8 | SNAP receptor | 0.2658 | 1.2 | 0.382028 |
| YIL046W | MET30 | protein binding | 0.2665 | 1.2 | 0.0675293 |
| YCL014W | BUD3 | unknown | 0.2708 | 1.2 | 0.376391 |
| YDL190C | UFD2 | ubiquitin conjugating enzyme | 0.2722 | 1.2 | 0.215452 |
| YLR293C | GSP1 | GTPase | 0.2725 | 1.2 | 0.245176 |
| YHR154W | RTT107 | unknown | 0.2728 | 1.2 | 0.224814 |
| YAL021C | CCR4 | 3'-5'-exoribonuclease | 0.2735 | 1.2 | 0.133492 |
| YER014W | HEM14 | protoporphyrinogen oxidase | 0.2743 | 1.2 | 0.0020775 |
| YLR185W | RPL37A | structural constituent of ribosome | 0.2750 | 1.2 | 0.0170647 |
| YIL077C | | unknown | 0.2767 | 1.2 | 0.15259 |
| YBR086C | IST2 | unknown | 0.2770 | 1.2 | 0.10021 |
| YAL048C | GEM1 | GTPase | 0.2782 | 1.2 | 0.0444376 |
| YPL090C | RPS6A | structural constituent of ribosome | 0.2783 | 1.2 | 0.0693926 |
| YCR062W | | | 0.2785 | 1.2 | 0.051378 |
| YMR070W | MOT3 | DNA binding* | 0.2793 | 1.2 | 0.341216 |
| YDR210C-D | | | 0.2807 | 1.2 | 0.0832661 |
| YKL172W | EBP2 | unknown | 0.2808 | 1.2 | 0.220731 |
| YBR012W-A | | | 0.2832 | 1.2 | 0.0769723 |
| YLR410W | VIP1 | unknown | 0.2852 | 1.2 | 0.27802 |
| YIL118W | RHO3 | GTPase * | 0.2860 | 1.2 | 0.0241781 |
| YIL043C | CBR1 | cytochrome-b5 reductase | 0.2865 | 1.2 | 0.105024 |
| YLR340W | RPP0 | structural constituent of ribosome | 0.2867 | 1.2 | 0.0219009 |
| YGL031C | RPL24A | structural constituent of ribosome* | 0.2887 | 1.2 | 0.186987 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YJR028W | | | 0.2888 | 1.2 | 0.284732 |
| Q0060 | AI3 | endonuclease | 0.2902 | 1.2 | 0.0322297 |
| YPR065W | ROX1 | specific transcriptional repressor * | 0.2902 | 1.2 | 0.240371 |
| YGR148C | RPL24B | structural constituent of ribosome* | 0.2907 | 1.2 | 0.0228892 |
| YOL053C-A | | | 0.2907 | 1.2 | 0.001846 |
| YFL-TYB | | | 0.2913 | 1.2 | 0.0545275 |
| YIR023W | DAL81 | specific RNA polymerase II transcription factor | 0.2913 | 1.2 | 0.0761399 |
| YNL042W | BOP3 | unknown | 0.2917 | 1.2 | 0.0396361 |
| YOR043W | WHI2 | phosphatase activator | 0.2917 | 1.2 | 0.107319 |
| YOL098C | | metalloendopeptidase | 0.2920 | 1.2 | 0.023922 |
| YBL005W-A | | | 0.2927 | 1.2 | 0.0987231 |
| YBR214W | SDS24 | unknown | 0.2937 | 1.2 | 0.0616586 |
| YBR200W | BEM1 | protein binding | 0.2948 | 1.2 | 0.128751 |
| YBR112C | CYC8 | transcription coactivator * | 0.2985 | 1.2 | 0.164839 |
| YBR256C | RIB5 | riboflavin synthase | 0.2993 | 1.2 | 0.134346 |
| YER131W | RPS26B | structural constituent of ribosome | 0.3003 | 1.2 | 0.0131872 |
| YHR203C | RPS4B | structural constituent of ribosome | 0.3007 | 1.2 | 0.0361563 |
| YBR198C | TAF5 | general RNA polymerase II transcription factor | 0.3050 | 1.2 | 0.0776355 |
| YHR089C | GAR1 | RNA binding | 0.3062 | 1.2 | 0.148651 |
| YPL220W | RPL1A | structural constituent of ribosome | 0.3073 | 1.2 | 0.112991 |
| YGL150C | INO80 | ATPase * | 0.3080 | 1.2 | 0.0468705 |
| YGL150C | INO80 | ATPase * | 0.3080 | 1.2 | 0.0468705 |
| YDR533C | HSP31 | unfolded protein binding* | 0.3095 | 1.2 | 0.0030393 |
| YCR107W | AAD3 | aryl-alcohol dehydrogenase | 0.3100 | 1.2 | 0.160967 |
| YDR035W | ARO3 | 3-deoxy-7-phosphoheptulonate synthase | 0.3110 | 1.2 | 0.0920598 |
| YPL095C | EEB1 | unknown | 0.3110 | 1.2 | 0.491436 |
| YLR177W | | unknown | 0.3137 | 1.2 | 0.611505 |
| YGL197W | MDS3 | unknown | 0.3150 | 1.2 | 0.435973 |
| YMR124W | | unknown | 0.3170 | 1.2 | 0.0685866 |
| YBR213W | MET8 | ferrochelatase * | 0.3187 | 1.2 | 0.145926 |
| YAL060W | BDH1 | (R,R)-butanediol dehydrogenase | 0.3200 | 1.2 | 0.157399 |
| YOR293W | RPS10A | structural constituent of ribosome | 0.3218 | 1.2 | 0.300713 |
| YGL128C | CWC23 | unknown | 0.3223 | 1.3 | 0.0229767 |
| YMR051C | | | 0.3235 | 1.3 | 0.217876 |
| YNL162W | RPL42A | structural constituent of ribosome | 0.3235 | 1.3 | 0.505042 |
| YBR129C | OPY1 | unknown | 0.3237 | 1.3 | 0.0533527 |
| YBL072C | RPS8A | structural constituent of ribosome | 0.3238 | 1.3 | 0.0011832 |
| YBL072C | RPS8A | structural constituent of ribosome | 0.3238 | 1.3 | 0.0011832 |
| YBL101C | ECM21 | unknown | 0.3250 | 1.3 | 0.12519 |
| YBR229C | ROT2 | alpha-glucosidase | 0.3270 | 1.3 | 0.450772 |
| YBR108W | | unknown | 0.3278 | 1.3 | 0.0567532 |
| YNL241C | ZWF1 | glucose-6-phosphate 1-dehydrogenase | 0.3280 | 1.3 | 0.0649378 |
| YDR444W | | unknown | 0.3292 | 1.3 | 0.133095 |
| YBR203W | COS111 | unknown | 0.3293 | 1.3 | 0.116873 |
| YBR295W | PCA1 | ATPase, coupled to transmembrane movement of ions, phosphorylation | 0.3323 | 1.3 | 0.237237 |
| YKL035W | UGP1 | UTP:glucose-1-phosphate uridylyltransferase | 0.3323 | 1.3 | 0.128612 |
| YBR197C | | unknown | 0.3328 | 1.3 | 0.0120974 |
| YOR359W | VTS1 | RNA binding* | 0.3328 | 1.3 | 0.031499 |
| YCR038C | BUD5 | signal transducer * | 0.3338 | 1.3 | 0.333285 |
| YNL027W | CRZ1 | transcription factor | 0.3347 | 1.3 | 0.0598415 |
| YIL060W | | unknown | 0.3360 | 1.3 | 0.0623404 |
| YMR125W | STO1 | mRNA binding | 0.3362 | 1.3 | 0.0148437 |
| YCL020W | | | 0.3372 | 1.3 | 0.273203 |
| Q0140 | VAR1 | structural constituent of ribosome | 0.3377 | 1.3 | 0.0330321 |
| YKL180W | RPL17A | structural constituent of ribosome | 0.3397 | 1.3 | 0.0045244 |
| YKL094W | YJU3 | serine hydrolase | 0.3408 | 1.3 | 0.0042445 |
| YKL094W | YJU3 | serine hydrolase | 0.3408 | 1.3 | 0.0042445 |
| YBR113W | | | 0.3417 | 1.3 | 0.0155249 |
| YML113W | DAT1 | AT DNA binding | 0.3417 | 1.3 | 0.0977873 |
| YBR302C | COS2 | unknown | 0.3418 | 1.3 | 0.105539 |
| YGL122C | NAB2 | poly(A) binding | 0.3423 | 1.3 | 0.0495587 |
| YBR089W | | | 0.3485 | 1.3 | 0.167728 |
| YOR234C | RPL33B | structural constituent of ribosome | 0.3503 | 1.3 | 0.317068 |
| YBR222C | PCS60 | AMP binding | 0.3507 | 1.3 | 0.0094451 |
| YGL189C | RPS26A | structural constituent of ribosome | 0.3522 | 1.3 | 0.0073061 |
| YCR029C | | | 0.3535 | 1.3 | 0.0147998 |
| YIL108W | | metalloendopeptidase | 0.3567 | 1.3 | 0.0009852 |
| YIL108W | | metalloendopeptidase | 0.3567 | 1.3 | 0.0009852 |
| YBR128C | ATG14 | unknown | 0.3583 | 1.3 | 0.0955886 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| Q0255 | | unknown | 0.3588 | 1.3 | 0.0691292 |
| YEL060C | PRB1 | serine-type endopeptidase | 0.3597 | 1.3 | 0.13995 |
| YAL056W | GPB2 | signal transducer | 0.3600 | 1.3 | 0.239646 |
| YLR333C | RPS25B | structural constituent of ribosome | 0.3607 | 1.3 | 0.0061303 |
| YOR306C | MCH5 | transporter | 0.3627 | 1.3 | 0.0062809 |
| YLR448W | RPL6B | structural constituent of ribosome* | 0.3643 | 1.3 | 0.0996407 |
| YOR312C | RPL20B | structural constituent of ribosome | 0.3650 | 1.3 | 0.445677 |
| YBR049C | REB1 | RNA polymerase II transcription factor | 0.3665 | 1.3 | 0.0020136 |
| YNR052C | POP2 | 3'-5'-exoribonuclease | 0.3668 | 1.3 | 0.0132947 |
| YPL060W | LPE10 | magnesium ion transporter | 0.3678 | 1.3 | 0.0132464 |
| YPL060W | LPE10 | magnesium ion transporter | 0.3678 | 1.3 | 0.0132464 |
| YHR214C-B | | | 0.3710 | 1.3 | 0.0597427 |
| YDL227C | HO | endonuclease | 0.3717 | 1.3 | 0.0209921 |
| YOL143C | RIB4 | 6,7-dimethyl-8-ribityllumazine synthase | 0.3738 | 1.3 | 0.0426318 |
| YJL076W | NET1 | rDNA binding | 0.3743 | 1.3 | 0.201175 |
| YMR264W | CUE1 | protein binding | 0.3745 | 1.3 | 0.0838317 |
| YKR075C | | unknown | 0.3748 | 1.3 | 0.308079 |
| YLR383W | SMC6 | unknown | 0.3752 | 1.3 | 0.0595504 |
| YJL136C | RPS21B | structural constituent of ribosome | 0.3757 | 1.3 | 0.0924836 |
| YGR098C | ESP1 | cysteine-type endopeptidase | 0.3767 | 1.3 | 0.522543 |
| YNL180C | RHO5 | GTPase | 0.3772 | 1.3 | 0.0093627 |
| YNL180C | RHO5 | GTPase | 0.3772 | 1.3 | 0.0093627 |
| YIL153W | RRD1 | protein phosphatase type 2A regulator | 0.3782 | 1.3 | 0.0362004 |
| YIL153W | RRD1 | protein phosphatase type 2A regulator | 0.3782 | 1.3 | 0.0362004 |
| YBR075W | | | 0.3797 | 1.3 | 0.0320617 |
| YDL046W | NPC2 | unknown | 0.3798 | 1.3 | 0.0016228 |
| YAR019C | CDC15 | protein kinase | 0.3803 | 1.3 | 0.109336 |
| YAR027W | UIP3 | unknown | 0.3817 | 1.3 | 0.0337224 |
| YPL229W | | unknown | 0.3832 | 1.3 | 0.022244 |
| YBR172C | SMY2 | unknown | 0.3845 | 1.3 | 0.0882636 |
| YPL081W | RPS9A | structural constituent of ribosome | 0.3850 | 1.3 | 0.0892798 |
| YBR166C | TYR1 | prephenate dehydrogenase | 0.3855 | 1.3 | 0.0407031 |
| YHR202W | | unknown | 0.3862 | 1.3 | 0.111643 |
| YJL177W | RPL17B | structural constituent of ribosome | 0.3878 | 1.3 | 0.0124636 |
| YCR084C | TUP1 | general transcriptional repressor | 0.3892 | 1.3 | 0.0188019 |
| YDR500C | RPL37B | structural constituent of ribosome | 0.3898 | 1.3 | 0.107413 |
| YML042W | CAT2 | carnitine O-acetyltransferase | 0.3918 | 1.3 | 0.0578596 |
| YBR249C | ARO4 | 3-deoxy-7-phosphoheptulonate synthase | 0.3933 | 1.3 | 0.0183409 |
| YEL054C | RPL12A | structural constituent of ribosome | 0.3938 | 1.3 | 0.403793 |
| YPL238C | | | 0.3992 | 1.3 | 0.0006738 |
| YPL238C | | | 0.3992 | 1.3 | 0.0006738 |
| YBL027W | RPL19B | structural constituent of ribosome | 0.3993 | 1.3 | 0.0243631 |
| YCR052W | RSC6 | unknown | 0.4013 | 1.3 | 0.0364 |
| YER088C | DOT6 | unknown | 0.4020 | 1.3 | 0.125387 |
| YBR138C | | unknown | 0.4030 | 1.3 | 0.120619 |
| YDR432W | NPL3 | mRNA binding* | 0.4045 | 1.3 | 0.0421114 |
| YCR039C | MATALPHA2 | transcription corepressor | 0.4047 | 1.3 | 0.0600152 |
| YJL015C | | | 0.4057 | 1.3 | 0.0009602 |
| YBR110W | ALG1 | beta-1,4-mannosyltransferase | 0.4103 | 1.3 | 0.0222501 |
| YER074W | RPS24A | structural constituent of ribosome | 0.4130 | 1.3 | 0.120579 |
| YDL085W | NDE2 | NADH dehydrogenase | 0.4135 | 1.3 | 0.0352389 |
| YHR142W | CHS7 | unknown | 0.4155 | 1.3 | 0.0808834 |
| YJL141C | YAK1 | protein kinase | 0.4157 | 1.3 | 0.100752 |
| YHR152W | SPO12 | unknown | 0.4168 | 1.3 | 0.326566 |
| YNL069C | RPL16B | structural constituent of ribosome* | 0.4210 | 1.3 | 0.0873251 |
| YPR043W | RPL43A | structural constituent of ribosome | 0.4213 | 1.3 | 0.0960937 |
| YKL006W | RPL14A | structural constituent of ribosome* | 0.4235 | 1.3 | 0.12409 |
| YOR377W | ATF1 | alcohol O-acetyltransferase | 0.4262 | 1.3 | 0.133978 |
| YJL165C | HAL5 | protein kinase | 0.4272 | 1.3 | 0.0223582 |
| YER182W | FMP10 | unknown | 0.4277 | 1.3 | 0.0383488 |
| YOL144W | NOP8 | unknown | 0.4278 | 1.3 | 0.0647453 |
| YBL064C | PRX1 | thioredoxin peroxidase | 0.4280 | 1.3 | 0.0402983 |
| YHR149C | SKG6 | unknown | 0.4280 | 1.3 | 0.0062213 |
| YJR086W | STE18 | GTPase | 0.4285 | 1.3 | 0.115272 |
| YCL011C | GBP2 | RNA binding* | 0.4298 | 1.3 | 0.0009988 |
| YLR330W | CHS5 | unknown | 0.4302 | 1.3 | 0.101969 |
| YIR017C | MET28 | DNA binding* | 0.4327 | 1.3 | 0.0513199 |
| YER056C | FCY2 | cytosine-purine permease | 0.4358 | 1.4 | 0.0434912 |
| YLR035C-A | | | 0.4382 | 1.4 | 0.013454 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YBR187W | | unknown | 0.4395 | 1.4 | 0.0110898 |
| YDL222C | FMP45 | unknown | 0.4407 | 1.4 | 0.246699 |
| YIL055C | | unknown | 0.4428 | 1.4 | 0.0382352 |
| YCR096C | HMRA2 | unknown | 0.4430 | 1.4 | 0.0433916 |
| YIR006C | PAN1 | protein binding, bridging | 0.4467 | 1.4 | 0.0071099 |
| YOR173W | DCS2 | unknown | 0.4503 | 1.4 | 0.0080365 |
| YJL139C | YUR1 | mannosyltransferase | 0.4505 | 1.4 | 0.0473155 |
| YHR165C | PRP8 | RNA splicing factor | 0.4522 | 1.4 | 0.0734453 |
| YPL237W | SUI3 | translation initiation factor | 0.4552 | 1.4 | 0.0786988 |
| YHL033C | RPL8A | structural constituent of ribosome | 0.4570 | 1.4 | 0.199437 |
| YCL042W | | unknown | 0.4598 | 1.4 | 0.289034 |
| YHR145C | | | 0.4600 | 1.4 | 0.0079284 |
| YML024W | RPS17A | structural constituent of ribosome | 0.4615 | 1.4 | 0.0030291 |
| YCL067C | HMLALPHA2 | transcription corepressor | 0.4633 | 1.4 | 0.0076669 |
| YGR019W | UGA1 | 4-aminobutyrate transaminase | 0.4637 | 1.4 | 0.0107372 |
| YPL225W | | unknown | 0.4642 | 1.4 | 0.130315 |
| YFL020C | PAU5 | unknown | 0.4672 | 1.4 | 0.0480115 |
| YMR251W-A | HOR7 | unknown | 0.4703 | 1.4 | 0.0005489 |
| YDR002W | YRB1 | Ran GTPase binding | 0.4705 | 1.4 | 0.31041 |
| Q0070 | AI5 ALPHA | endonuclease | 0.4720 | 1.4 | 0.0212955 |
| Q0070 | AI5 ALPHA | endonuclease | 0.4720 | 1.4 | 0.0212955 |
| YPL004C | LSP1 | protein kinase | 0.4748 | 1.4 | 0.009949 |
| YDR072C | IPT1 | transferase , transferring phosphorus-containing groups | 0.4757 | 1.4 | 0.176764 |
| YJR026W | | | 0.4777 | 1.4 | 0.165214 |
| YHR134W | WSS1 | unknown | 0.4810 | 1.4 | 0.0067361 |
| YPR102C | RPL11A | structural constituent of ribosome | 0.4812 | 1.4 | 0.0245074 |
| YDL075W | RPL31A | structural constituent of ribosome | 0.4855 | 1.4 | 0.0346551 |
| YDL142C | CRD1 | cardiolipin synthase | 0.4862 | 1.4 | 0.0005569 |
| YDL142C | CRD1 | cardiolipin synthase | 0.4862 | 1.4 | 0.0005569 |
| YDL232W | OST4 | protein binding, bridging* | 0.4865 | 1.4 | 0.0183406 |
| YDL088C | ASM4 | structural molecule | 0.4878 | 1.4 | 0.167887 |
| YMR194W | RPL36A | structural constituent of ribosome* | 0.4903 | 1.4 | 0.0085836 |
| YIR017W-A | | | 0.4927 | 1.4 | 0.115578 |
| YBR048W | RPS11B | structural constituent of ribosome | 0.4943 | 1.4 | 0.0705719 |
| YBR122C | MRPL36 | structural constituent of ribosome | 0.4948 | 1.4 | 0.163655 |
| YBR062C | | unknown | 0.4957 | 1.4 | 0.0397618 |
| YCR031 | | | 0.4963 | 1.4 | 0.0853844 |
| YCL024W | KCC4 | protein kinase | 0.4987 | 1.4 | 0.0728841 |
| YJL118W | | unknown | 0.4987 | 1.4 | 0.124295 |
| YCR045C | | serine-type peptidase | 0.5000 | 1.4 | 0.169024 |
| YDL191W | RPL35A | structural constituent of ribosome | 0.5005 | 1.4 | 0.103185 |
| YLR414C | | unknown | 0.5023 | 1.4 | 0.0111992 |
| YNL284C-A | | | 0.5025 | 1.4 | 0.0549954 |
| YAR009C | | | 0.5025 | 1.4 | 0.0040989 |
| YML063W | RPS1B | structural constituent of ribosome | 0.5028 | 1.4 | 0.004124 |
| YJL065C | DLS1 | unknown | 0.5045 | 1.4 | 0.0538968 |
| YIR012W | SQT1 | unknown | 0.5067 | 1.4 | 0.100443 |
| YOL127W | RPL25 | structural constituent of ribosome* | 0.5103 | 1.4 | 0.12283 |
| YIL074C | SER33 | phosphoglycerate dehydrogenase | 0.5158 | 1.4 | 0.010749 |
| YOL040C | RPS15 | structural constituent of ribosome | 0.5160 | 1.4 | 0.273104 |
| YDL188C | PPH2 | protein phosphatase type 2A | 0.5198 | 1.4 | 0.0550047 |
| YLR237W | THI7 | thiamin transporter | 0.5210 | 1.4 | 0.0427702 |
| YGL078C | DBP3 | ATP-dependent RNA helicase | 0.5217 | 1.4 | 0.15154 |
| YIL173W | VTH1 | signal sequence binding | 0.5225 | 1.4 | 0.244468 |
| YLR344W | RPL26A | structural constituent of ribosome* | 0.5235 | 1.4 | 0.0825584 |
| YDL053C | PBP4 | unknown | 0.5237 | 1.4 | 0.0990367 |
| YER102W | RPS8B | structural constituent of ribosome | 0.5258 | 1.4 | 0.005912 |
| YBR084C-A | RPL19A | structural constituent of ribosome | 0.5270 | 1.4 | 0.0244341 |
| YML073C | RPL6A | structural constituent of ribosome* | 0.5292 | 1.4 | 0.0214537 |
| YDR041W | RSM10 | structural constituent of ribosome | 0.5327 | 1.4 | 0.0096728 |
| YDL243C | AAD4 | aryl-alcohol dehydrogenase | 0.5348 | 1.4 | 0.0060642 |
| YDL082W | RPL13A | structural constituent of ribosome | 0.5392 | 1.5 | 0.256101 |
| YOR052C | | unknown | 0.5397 | 1.5 | 0.137075 |
| YDL136W | RPL35B | structural constituent of ribosome | 0.5422 | 1.5 | 0.125916 |
| YGR137W | | | 0.5445 | 1.5 | 8.151E-05 |
| YNL244C | SUI1 | translation initiation factor | 0.5460 | 1.5 | 0.0053681 |
| YGL135W | RPL1B | structural constituent of ribosome | 0.5503 | 1.5 | 0.0087886 |
| YER159C | BUR6 | transcription corepressor | 0.5523 | 1.5 | 0.0147145 |
| YDL020C | RPN4 | transcriptional activator | 0.5535 | 1.5 | 0.0598933 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YCR024C-A | PMP1 | enzyme regulator | 0.5552 | 1.5 | 0.0023886 |
| YDR261C-C | | | 0.5552 | 1.5 | 0.0020688 |
| YCR008W | SAT4 | protein kinase | 0.5563 | 1.5 | 0.0170853 |
| YDR210W-A | | | 0.5570 | 1.5 | 0.186916 |
| YLR384C | IKI3 | RNA polymerase II transcription elongation factor | 0.5592 | 1.5 | 0.0575922 |
| YDR261W-A | | | 0.5607 | 1.5 | 0.0048746 |
| YHR010W | RPL27A | structural constituent of ribosome | 0.5617 | 1.5 | 0.0254427 |
| YDR447C | RPS17B | structural constituent of ribosome | 0.5633 | 1.5 | 0.295822 |
| YLR441C | RPS1A | structural constituent of ribosome | 0.5658 | 1.5 | 0.076886 |
| YHR155W | YSP1 | unknown | 0.5687 | 1.5 | 0.0970743 |
| YBR068C | BAP2 | amino acid transporter | 0.5702 | 1.5 | 0.0664958 |
| YAR031W | PRM9 | unknown | 0.5723 | 1.5 | 0.219092 |
| YLR250W | SSP120 | unknown | 0.5730 | 1.5 | 0.0039065 |
| YJL117W | PHO86 | unknown | 0.5767 | 1.5 | 0.0304931 |
| YOR122C | PFY1 | actin monomer binding | 0.5872 | 1.5 | 0.109343 |
| YPL143W | RPL33A | structural constituent of ribosome | 0.5910 | 1.5 | 0.0781045 |
| YPL131W | RPL5 | structural constituent of ribosome* | 0.5928 | 1.5 | 0.0278664 |
| YHR141C | RPL42B | structural constituent of ribosome | 0.5947 | 1.5 | 0.0040292 |
| YHL015W | RPS20 | structural constituent of ribosome | 0.6025 | 1.5 | 0.0375388 |
| YEL046C | GLY1 | threonine aldolase | 0.6060 | 1.5 | 0.0162237 |
| YER064C | | unknown | 0.6093 | 1.5 | 0.114616 |
| YIL069C | RPS24B | structural constituent of ribosome | 0.6097 | 1.5 | 0.0138118 |
| YDR074W | TPS2 | trehalose-phosphatase | 0.6127 | 1.5 | 0.10452 |
| YHR097C | | unknown | 0.6127 | 1.5 | 0.0033287 |
| YDR365W-A | | | 0.6128 | 1.5 | 0.0082528 |
| YBR154C | RPB5 | DNA-directed RNA polymerase | 0.6132 | 1.5 | 0.283729 |
| YJL122W | ALB1 | unknown | 0.6147 | 1.5 | 0.136374 |
| YGL095C | VPS45 | unfolded protein binding | 0.6153 | 1.5 | 0.0130891 |
| YLR382C | NAM2 | mRNA binding* | 0.6177 | 1.5 | 0.0317762 |
| YBR183W | YPC1 | ceramidase | 0.6182 | 1.5 | 0.0486957 |
| YMR143W | RPS16A | structural constituent of ribosome | 0.6187 | 1.5 | 0.0354319 |
| YDL048C | STP4 | unknown | 0.6193 | 1.5 | 0.0489642 |
| YMR107W | SPG4 | unknown | 0.6218 | 1.5 | 0.15285 |
| YDR418W | RPL12B | structural constituent of ribosome | 0.6225 | 1.5 | 0.0178555 |
| YMR142C | RPL13B | structural constituent of ribosome | 0.6235 | 1.5 | 0.0087378 |
| YKL096W | CWP1 | structural constituent of cell wall | 0.6240 | 1.5 | 0.0539334 |
| YDR366C | | unknown | 0.6243 | 1.5 | 0.0298718 |
| YNL067W | RPL9B | structural constituent of ribosome | 0.6250 | 1.5 | 0.0875433 |
| YOR192C-A | | | 0.6257 | 1.5 | 0.0534534 |
| YJL191W | RPS14B | structural constituent of ribosome* | 0.6272 | 1.5 | 0.0040079 |
| YJL189W | RPL39 | structural constituent of ribosome | 0.6295 | 1.5 | 0.0222771 |
| YNL268W | LYP1 | basic amino acid transporter | 0.6315 | 1.5 | 0.0568801 |
| YGL077C | HNM1 | choline transporter | 0.6335 | 1.6 | 0.355304 |
| YLR107W | REX3 | 3'-5' exonuclease | 0.6347 | 1.6 | 0.134479 |
| YLR388W | RPS29A | structural constituent of ribosome | 0.6367 | 1.6 | 0.0117181 |
| YPL061W | ALD6 | 3-chloroallyl aldehyde dehydrogenase | 0.6368 | 1.6 | 0.0187111 |
| YDR269C | | | 0.6383 | 1.6 | 0.0001662 |
| YBL035C | POL12 | alpha DNA polymerase | 0.6437 | 1.6 | 0.0223498 |
| YBR190W | | | 0.6440 | 1.6 | 0.0243461 |
| YER049W | TPA1 | unknown | 0.6458 | 1.6 | 0.0726418 |
| YBR114W | RAD16 | DNA-dependent ATPase | 0.6507 | 1.6 | 0.0018041 |
| YDL061C | RPS29B | structural constituent of ribosome | 0.6518 | 1.6 | 0.0702054 |
| YIL152W | | unknown | 0.6540 | 1.6 | 0.0139991 |
| YOR309C | | | 0.6545 | 1.6 | 0.0631489 |
| YCR050C | | | 0.6575 | 1.6 | 0.0312211 |
| YJL192C | SOP4 | unknown | 0.6612 | 1.6 | 0.0380758 |
| YJL192C | SOP4 | unknown | 0.6612 | 1.6 | 0.0380758 |
| YCL040W | GLK1 | glucokinase | 0.6623 | 1.6 | 0.0435096 |
| YDR034C-C | | | 0.6717 | 1.6 | 0.010173 |
| YOR080W | DIA2 | DNA replication origin binding | 0.6753 | 1.6 | 0.0061567 |
| YHL002W | HSE1 | protein binding | 0.6760 | 1.6 | 0.0183859 |
| YHR166C | CDC23 | protein binding* | 0.6782 | 1.6 | 0.0133522 |
| YHR187W | IKI1 | RNA polymerase II transcription elongation factor | 0.6812 | 1.6 | 0.0506781 |
| YGR085C | RPL11B | structural constituent of ribosome | 0.6815 | 1.6 | 0.0106195 |
| YJL144W | | unknown | 0.6815 | 1.6 | 0.146826 |
| YBR191W | RPL21A | structural constituent of ribosome | 0.6820 | 1.6 | 0.0224524 |
| YKL065C | YET1 | unknown | 0.6908 | 1.6 | 0.0149104 |
| YHL001W | RPL14B | structural constituent of ribosome* | 0.6952 | 1.6 | 0.0124379 |
| YCRX16C | | | 0.6958 | 1.6 | 0.0465628 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YDR025W | RPS11A | structural constituent of ribosome | 0.6967 | 1.6 | 0.0171773 |
| YIL144W | TID3 | structural constituent of cytoskeleton | 0.7028 | 1.6 | 0.0248803 |
| YMR170C | ALD2 | 3-chloroalyl aldehyde dehydrogenase | 0.7032 | 1.6 | 0.116715 |
| YKR097W | PCK1 | phosphoenolpyruvate carboxykinase | 0.7040 | 1.6 | 0.0130773 |
| YGL123W | RPS2 | structural constituent of ribosome | 0.7088 | 1.6 | 0.0029485 |
| YDL077C | VAM6 | Rab guanyl-nucleotide exchange factor | 0.7118 | 1.6 | 0.0281424 |
| YKL006C-A | SFT1 | v-SNARE | 0.7140 | 1.6 | 0.0118243 |
| YDL083C | RPS16B | structural constituent of ribosome | 0.7155 | 1.6 | 0.0033495 |
| YJL089W | SIP4 | specific RNA polymerase II transcription factor | 0.7180 | 1.6 | 0.0646857 |
| YER056C-A | RPL34A | structural constituent of ribosome | 0.7208 | 1.6 | 0.0106918 |
| YIR031C | DAL7 | malate synthase | 0.7313 | 1.7 | 0.0109947 |
| YNL054W-A | | | 0.7348 | 1.7 | 0.0386707 |
| YIL052C | RPL34B | structural constituent of ribosome | 0.7393 | 1.7 | 0.0467228 |
| YHR138C | | endopeptidase inhibitor | 0.7437 | 1.7 | 0.0011383 |
| YLR410W-A | | | 0.7458 | 1.7 | 0.0009951 |
| YBR188C | NTC20 | RNA splicing factor | 0.7475 | 1.7 | 0.0117737 |
| YJR123W | RPS5 | structural constituent of ribosome | 0.7492 | 1.7 | 0.0536007 |
| YJL171C | | unknown | 0.7520 | 1.7 | 0.0277478 |
| YKL156W | RPS27A | structural constituent of ribosome | 0.7547 | 1.7 | 0.0727957 |
| YGR136W | LSB1 | unknown | 0.7552 | 1.7 | 0.0254165 |
| YKL024C | URA6 | uridine kinase | 0.7587 | 1.7 | 0.0737835 |
| YBL092W | RPL32 | structural constituent of ribosome | 0.7623 | 1.7 | 0.0333798 |
| YLL045C | RPL8B | structural constituent of ribosome | 0.7663 | 1.7 | 0.0074899 |
| YFL031W | HAC1 | transcription factor | 0.7707 | 1.7 | 0.0417002 |
| YHR021C | RPS27B | structural constituent of ribosome | 0.7717 | 1.7 | 0.0141383 |
| YAR037W | | | 0.7727 | 1.7 | 0.0830785 |
| YNL113W | RPC19 | DNA-directed RNA polymerase | 0.7785 | 1.7 | 0.0580063 |
| YDL051W | LHP1 | RNA binding | 0.7820 | 1.7 | 0.0227754 |
| YLL039C | UBI4 | protein tag* | 0.7848 | 1.7 | 0.0007606 |
| YDR203W | | | 0.7855 | 1.7 | 0.0129925 |
| YPL079W | RPL21B | structural constituent of ribosome | 0.7868 | 1.7 | 0.0651095 |
| YJR095W | SFC1 | succinate:fumarate antiporter | 0.7895 | 1.7 | 0.0225915 |
| YMR316W | DIA1 | unknown | 0.7928 | 1.7 | 0.0027432 |
| YDL134C | PPH21 | protein phosphatase type 2A | 0.7975 | 1.7 | 0.013999 |
| YJL190C | RPS22A | structural constituent of ribosome | 0.7998 | 1.7 | 0.0013096 |
| YJR096W | | aldo-keto reductase * | 0.8017 | 1.7 | 0.121383 |
| YHR095W | | | 0.8072 | 1.7 | 0.0694593 |
| YMR242C | RPL20A | structural constituent of ribosome | 0.8078 | 1.8 | 0.0002689 |
| YMR243C | ZRC1 | di-, tri-valent inorganic cation transporter | 0.8082 | 1.8 | 0.117849 |
| YLR076C | | | 0.8155 | 1.8 | 0.016312 |
| YMR191W | SPG5 | unknown | 0.8167 | 1.8 | 0.0200889 |
| YLR267W | BOP2 | unknown | 0.8172 | 1.8 | 0.015504 |
| YLR267W | BOP2 | unknown | 0.8172 | 1.8 | 0.015504 |
| YDL170W | UGA3 | transcription factor | 0.8173 | 1.8 | 0.0202773 |
| YCR035C | RRP43 | 3'-5'-exoribonuclease | 0.8242 | 1.8 | 0.0046852 |
| YML026C | RPS18B | structural constituent of ribosome | 0.8287 | 1.8 | 0.131321 |
| YDR471W | RPL27B | structural constituent of ribosome | 0.8310 | 1.8 | 0.0159222 |
| YDL134C-A | | | 0.8322 | 1.8 | 0.0421843 |
| YDR450W | RPS18A | structural constituent of ribosome | 0.8353 | 1.8 | 0.0071381 |
| YDR055W | PST1 | unknown | 0.8372 | 1.8 | 0.0160348 |
| YEL026W | SNU13 | RNA binding* | 0.8375 | 1.8 | 0.0614777 |
| YKL151C | | unknown | 0.8380 | 1.8 | 0.0009637 |
| YER013W | PRP22 | RNA splicing factor | 0.8392 | 1.8 | 0.0037659 |
| YIR033W | MGA2 | transcriptional activator | 0.8393 | 1.8 | 0.0177476 |
| YDL110C | TMA17 | unknown | 0.8412 | 1.8 | 0.0084966 |
| YHR137W | ARO9 | aromatic-amino-acid transaminase | 0.8435 | 1.8 | 0.01094 |
| YCR041W | | | 0.8443 | 1.8 | 0.0286948 |
| YMR176W | ECM5 | unknown | 0.8462 | 1.8 | 0.0143677 |
| YMR116C | ASC1 | unknown | 0.8542 | 1.8 | 0.0607158 |
| YCR033W | SNT1 | NAD-dependent histone deacetylase * | 0.8575 | 1.8 | 0.0187163 |
| YHR199C | FMP34 | unknown | 0.8585 | 1.8 | 0.01634 |
| YOR096W | RPS7A | structural constituent of ribosome | 0.8640 | 1.8 | 0.0035532 |
| YDL047W | SIT4 | protein serine/threonine phosphatase | 0.8682 | 1.8 | 0.0194416 |
| YBR004C | GPI18 | mannosyltransferase | 0.8723 | 1.8 | 0.0351371 |
| YBR117C | TKL2 | transketolase | 0.8727 | 1.8 | 0.0238077 |
| YBR116C | | | 0.8772 | 1.8 | 0.0255202 |
| YPL057C | SUR1 | mannosyltransferase * | 0.8797 | 1.8 | 0.0398194 |
| YNL192W | CHS1 | chitin synthase | 0.8813 | 1.8 | 0.0507604 |
| YGR086C | PIL1 | protein kinase | 0.8832 | 1.8 | 0.0068342 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YKR040C | | | 0.8837 | 1.8 | 0.0099214 |
| YML128C | MSC1 | unknown | 0.8883 | 1.9 | 0.0095036 |
| YDL204W | RTN2 | unknown | 0.8890 | 1.9 | 0.0026248 |
| YOR316C | COT1 | di-, tri-valent inorganic cation transporter | 0.8907 | 1.9 | 0.0109291 |
| YIL133C | RPL16A | structural constituent of ribosome* | 0.8953 | 1.9 | 0.0154846 |
| YEL074W | | | 0.8975 | 1.9 | 0.0178998 |
| YNL096C | RPS7B | structural constituent of ribosome | 0.8980 | 1.9 | 0.0790623 |
| YCR028C | FEN2 | pantothenate transporter | 0.9075 | 1.9 | 0.0068358 |
| YGL103W | RPL28 | structural constituent of ribosome* | 0.9087 | 1.9 | 0.0103456 |
| YOR095C | RKI1 | ribose-5-phosphate isomerase | 0.9110 | 1.9 | 0.0153279 |
| YNL117W | MLS1 | malate synthase | 0.9125 | 1.9 | 0.0173036 |
| YLR075W | RPL10 | structural constituent of ribosome | 0.9155 | 1.9 | 0.042146 |
| YDL212W | SHR3 | unfolded protein binding | 0.9160 | 1.9 | 0.0024472 |
| YFL050C | ALR2 | di-, tri-valent inorganic cation transporter | 0.9200 | 1.9 | 0.0048175 |
| YNL157W | IGO1 | unknown | 0.9290 | 1.9 | 0.0123405 |
| YBR031W | RPL4A | structural constituent of ribosome | 0.9317 | 1.9 | 0.0108182 |
| YBR050C | REG2 | protein phosphatase type 1 regulator | 0.9388 | 1.9 | 0.008376 |
| YBR297W | MAL33 | transcription factor | 0.9447 | 1.9 | 0.0159806 |
| YIL148W | RPL40A | structural constituent of ribosome* | 0.9465 | 1.9 | 0.0067275 |
| YOL126C | MDH2 | L-malate dehydrogenase | 0.9487 | 1.9 | 0.0028429 |
| YMR262W | | unknown | 0.9533 | 1.9 | 0.0271089 |
| YLR375W | STP3 | unknown | 0.9573 | 1.9 | 0.0050155 |
| YDL182W | LYS20 | homocitrate synthase | 0.9578 | 1.9 | 0.0231322 |
| YCL039W | GID7 | unknown | 0.9585 | 1.9 | 0.0417765 |
| YGR034W | RPL26B | structural constituent of ribosome* | 0.9658 | 2.0 | 0.0021637 |
| YLR029C | RPL15A | structural constituent of ribosome* | 0.9703 | 2.0 | 0.032712 |
| YDR001C | NTH1 | alpha, alpha-trehalase | 0.9763 | 2.0 | 0.0546775 |
| YDL123W | SNA4 | unknown | 0.9892 | 2.0 | 0.022495 |
| YNL195C | | unknown | 0.9907 | 2.0 | 0.0125122 |
| YGR088W | CTT1 | catalase | 0.9985 | 2.0 | 0.0019508 |
| YBR033W | EDS1 | unknown | 1.0075 | 2.0 | 0.111556 |
| YDR368W | YPR1 | oxidoreductase * | 1.0172 | 2.0 | 0.0406334 |
| YMR272C | SCS7 | oxidoreductase | 1.0177 | 2.0 | 0.0009907 |
| YER081W | SER3 | phosphoglycerate dehydrogenase | 1.0217 | 2.0 | 0.0029483 |
| YLR335C | ILV5 | ketol-acid reductoisomerase | 1.0233 | 2.0 | 0.0031752 |
| YER025W | GCD11 | translation initiation factor | 1.0372 | 2.1 | 0.0159438 |
| YDR012W | RPL4B | structural constituent of ribosome | 1.0437 | 2.1 | 0.0033139 |
| YBR161W | CSH1 | mannosyltransferase * | 1.0477 | 2.1 | 0.0041389 |
| YNL143C | | | 1.0530 | 2.1 | 0.0145176 |
| YML004C | GLO1 | lactoylglutathione lyase | 1.0553 | 2.1 | 0.000554 |
| YBR034C | HMT1 | protein-arginine N-methyltransferase | 1.0607 | 2.1 | 0.0004575 |
| YMR174C | PAI3 | endopeptidase inhibitor | 1.0615 | 2.1 | 0.0139104 |
| YNL178W | RPS3 | structural constituent of ribosome | 1.0698 | 2.1 | 0.0004945 |
| YER150W | SPI1 | unknown | 1.0750 | 2.1 | 0.0338064 |
| YHR049W | FSH1 | serine hydrolase | 1.0812 | 2.1 | 0.0863549 |
| YHR071W | PCL5 | cyclin-dependent protein kinase | 1.0830 | 2.1 | 0.0369871 |
| YDR508C | GNP1 | amino acid transporter | 1.0887 | 2.1 | 0.0089435 |
| YDL023C | | | 1.0908 | 2.1 | 0.0343537 |
| YCR005C | CIT2 | citrate (Si)-synthase | 1.0922 | 2.1 | 0.0174329 |
| YOR063W | RPL3 | structural constituent of ribosome | 1.0960 | 2.1 | 0.0107875 |
| YMR175W | SIP18 | phospholipid binding | 1.0990 | 2.1 | 0.000566 |
| YDR070C | FMP16 | unknown | 1.1073 | 2.2 | 0.0165054 |
| YBL043W | ECM13 | unknown | 1.1287 | 2.2 | 0.0023883 |
| YDR064W | RPS13 | structural constituent of ribosome | 1.1408 | 2.2 | 0.0580586 |
| YGR043C | | transaldolase | 1.1483 | 2.2 | 0.0072384 |
| YBR189W | RPS9B | structural constituent of ribosome | 1.1552 | 2.2 | 0.0069421 |
| YHR161C | YAP1801 | clathrin binding | 1.1663 | 2.2 | 0.0023362 |
| YML070W | DAK1 | glycerone kinase | 1.1700 | 2.3 | 0.0455597 |
| YGR057C | LST7 | protein transporter | 1.1775 | 2.3 | 0.0033761 |
| YGR057C | LST7 | protein transporter | 1.1775 | 2.3 | 0.0033761 |
| YML131W | | unknown | 1.1815 | 2.3 | 0.0427881 |
| YML131W | | unknown | 1.1815 | 2.3 | 0.0427881 |
| YER065C | ICL1 | isocitrate lyase | 1.1993 | 2.3 | 0.0087595 |
| YBL093C | ROX3 | RNA polymerase II transcription mediator | 1.2070 | 2.3 | 0.0064967 |
| YBL093C | ROX3 | RNA polymerase II transcription mediator | 1.2070 | 2.3 | 0.0064967 |
| YGR138C | TPO2 | spermine transporter | 1.2137 | 2.3 | 0.0014605 |
| YGR051C | | | 1.2417 | 2.4 | 0.0067508 |
| YMR173W | DDR48 | ATPase * | 1.2450 | 2.4 | 0.0176233 |
| YDR037W | KRS1 | lysine-tRNA ligase | 1.2452 | 2.4 | 0.006399 |

Table 2: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YGL037C | PNC1 | nicotinamidase | 1.2557 | 2.4 | 0.0090876 |
| YHR171W | ATG7 | APG12 activating enzyme * | 1.2777 | 2.4 | 0.0007471 |
| YHR104W | GRE3 | aldo-keto reductase * | 1.3027 | 2.5 | 0.002315 |
| YMR173W-A | | | 1.3243 | 2.5 | 0.0064815 |
| YDR492W | IZH1 | metal ion binding | 1.3452 | 2.5 | 0.0159989 |
| YLR194C | | structural constituent of cell wall | 1.3477 | 2.5 | 0.0121587 |
| YLR194C | | structural constituent of cell wall | 1.3477 | 2.5 | 0.0121587 |
| YOL125W | | unknown | 1.3705 | 2.6 | 0.0233996 |
| YOL125W | | unknown | 1.3705 | 2.6 | 0.0233996 |
| YMR090W | | unknown | 1.3865 | 2.6 | 0.0319874 |
| YNL112W | DBP2 | RNA helicase | 1.3868 | 2.6 | 0.0633853 |
| YFL021W | GAT1 | specific RNA polymerase II transcription factor | 1.3957 | 2.6 | 0.0314206 |
| YPL223C | GRE1 | unknown | 1.4073 | 2.7 | 0.0008462 |
| YBL033C | RIB1 | cyclohydrolase | 1.4123 | 2.7 | 0.0519619 |
| YFL029C | CAK1 | cyclin-dependent protein kinase | 1.4128 | 2.7 | 0.0009785 |
| YER024W | YAT2 | carnitine O-acetyltransferase | 1.4378 | 2.7 | 0.0105631 |
| YBR105C | VID24 | unknown | 1.5025 | 2.8 | 0.0160432 |
| YNL036W | NCE103 | carbonate dehydratase | 1.5077 | 2.8 | 0.0046861 |
| YIL117C | PRM5 | unknown | 1.5107 | 2.8 | 0.0013873 |
| YIL117C | PRM5 | unknown | 1.5107 | 2.8 | 0.0013873 |
| YOR348C | PUT4 | L-proline permease | 1.5480 | 2.9 | 0.0034499 |
| YDR038C | ENA5 | ATPase , coupled to transmembrane movement of ions, phosphorylation | 1.5610 | 3.0 | 0.0144567 |
| YDR038C | ENA5 | ATPase , coupled to transmembrane movement of ions, phosphorylation | 1.5610 | 3.0 | 0.0144567 |
| YDR502C | SAM2 | methionine adenosyltransferase | 1.5610 | 3.0 | 0.0020633 |
| YMR195W | ICY1 | unknown | 1.6000 | 3.0 | 0.0062503 |
| YDL168W | SFA1 | alcohol dehydrogenase | 1.6098 | 3.1 | 0.0072298 |
| YHR054C | | unknown | 1.6337 | 3.1 | 0.042637 |
| YER032W | FIR1 | unknown | 1.6533 | 3.1 | 0.0069945 |
| YIL099W | SGA1 | glucan 1,4-alpha-glucosidase | 1.6637 | 3.2 | 0.0273178 |
| YCR010C | ADY2 | ammonium transporter | 1.6937 | 3.2 | 0.0198648 |
| YPL135W | ISU1 | protein binding | 1.7005 | 3.3 | 0.0026833 |
| YHR139C | SPS100 | unknown | 1.7403 | 3.3 | 0.015393 |
| YJL088W | ARG3 | ornithine carbamoyltransferase | 1.7542 | 3.4 | 0.0138 |
| YDR040C | ENA1 | ATPase , coupled to transmembrane movement of ions, phosphorylation | 1.7885 | 3.5 | 0.0563136 |
| YOR031W | CRS5 | copper ion binding | 1.9118 | 3.8 | 0.0165351 |
| YFL013W-A | | | 1.9137 | 3.8 | 0.0008942 |
| YEL071W | DLD3 | D-lactate dehydrogenase | 1.9187 | 3.8 | 0.0016436 |
| YFL014W | HSP12 | unknown | 1.9348 | 3.8 | 0.0012294 |
| YHR103W | SBE22 | unknown | 1.9427 | 3.8 | 0.0115856 |
| YDR039C | ENA2 | ATPase , coupled to transmembrane movement of ions, phosphorylation | 1.9688 | 3.9 | 0.005857 |
| YLR061W | RPL22A | structural constituent of ribosome | 2.0047 | 4.0 | 0.0090796 |
| YHR162W | | unknown | 2.0162 | 4.0 | 0.0030272 |
| YOR303W | CPA1 | carbamoyl-phosphate synthase (glutamine-hydrolyzing) | 2.0510 | 4.1 | 0.0016321 |
| YDL022W | GPD1 | glycerol-3-phosphate dehydrogenase | 2.0962 | 4.3 | 0.0007573 |
| YDR043C | NRG1 | DNA binding* | 2.1978 | 4.6 | 0.0213979 |
| YHR070W | TRM5 | tRNA (guanine) methyltransferase | 2.2173 | 4.7 | 0.0342632 |
| YER145C | FTR1 | iron ion transporter | 2.2430 | 4.7 | 0.0084955 |
| YCL065W | | | 2.2572 | 4.8 | 0.0163941 |
| YDL214C | PRR2 | receptor signaling protein serine/threonine kinase | 2.2830 | 4.9 | 0.0124674 |
| YHR033W | | unknown | 2.4160 | 5.3 | 0.0027482 |
| YHR056C | RSC30 | DNA binding | 2.4358 | 5.4 | 0.0027017 |
| YFL035C-B | | | 2.5072 | 5.7 | 0.0324437 |
| YHR087W | | unknown | 2.5665 | 5.9 | 0.0054202 |
| YGL039W | | oxidoreductase * | 2.6283 | 6.2 | 0.031805 |
| YOR100C | CRC1 | carnitine:acyl carnitine antiporter | 2.7913 | 6.9 | 0.0004714 |
| YOL150C | | | 2.8075 | 7.0 | 0.0182197 |
| YJL106W | IME2 | protein kinase | 3.1218 | 8.7 | 0.0085618 |
| YIL053W | RHR2 | glycerol-1-phosphatase | 3.1282 | 8.7 | 0.0009198 |
| YGL157W | | oxidoreductase * | 3.2108 | 9.3 | 0.0064748 |
| YER062C | HOR2 | glycerol-1-phosphatase | 3.2197 | 9.3 | 0.0005317 |
| YOR107W | RGS2 | GTPase activator | 3.2452 | 9.5 | 0.0051711 |
| YMR169C | ALD3 | 3-chloroallyl aldehyde dehydrogenase | 3.2817 | 9.7 | 0.0119141 |
| YBR281C | | unknown | 3.4813 | 11.2 | 0.0001763 |
| YDR536W | STL1 | transporter | 3.5778 | 11.9 | 0.0024304 |
| YHR055C | CUP1-2 | copper ion binding | 3.7490 | 13.4 | 0.0004694 |
| YOL151W | GRE2 | oxidoreductase * | 3.7803 | 13.7 | 0.0034423 |
| YHR053C | CUP1-1 | copper ion binding | 3.8430 | 14.4 | 0.0003756 |
| YGR159C | NSR1 | RNA binding* | 3.8437 | 14.4 | 0.0007395 |
| YGR066C | | unknown | 4.3562 | 20.5 | 0.0112158 |

Table 3: Raw microarray data for fermentation day four.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|------------|---|----------------------------|-------------|-----------|
| YGR254W | ENO1 | phosphopyruvate hydratase | -3.5377 | -11.6 | 0.0005211 |
| YPL281C | ERR2 | phosphopyruvate hydratase | -3.5108 | -11.4 | 5.267E-05 |
| YDR246W | TRS23 | unknown | -3.3958 | -10.5 | 0.0013327 |
| YFL011W | HXT10 | glucose transporter | -3.2197 | -9.3 | 0.0088991 |
| YDR247W | VHS1 | protein kinase | -3.1373 | -8.8 | 0.0029673 |
| YDR249C | | unknown | -2.8802 | -7.4 | 0.0049436 |
| YNL142W | MEP2 | ammonium transporter | -2.8245 | -7.1 | 0.0029756 |
| YHR174W | ENO2 | phosphopyruvate hydratase | -2.8163 | -7.0 | 0.016026 |
| YFL030W | AGX1 | alanine-glyoxylate transaminase | -2.7903 | -6.9 | 0.0019559 |
| YGR087C | PDC6 | pyruvate decarboxylase | -2.7440 | -6.7 | 0.0040176 |
| YHR173C | | | -2.7333 | -6.6 | 0.0685791 |
| YHR181W | SVP26 | protein binding | -2.7217 | -6.6 | 0.0053433 |
| YLR153C | ACS2 | acetate-CoA ligase | -2.6993 | -6.5 | 0.0523336 |
| YOL086C | ADH1 | alcohol dehydrogenase | -2.6982 | -6.5 | 7.147E-05 |
| YMR303C | ADH2 | alcohol dehydrogenase | -2.6880 | -6.4 | 0.0007619 |
| YMR323W | ERR3 | phosphopyruvate hydratase | -2.6003 | -6.1 | 0.0160369 |
| YCR012W | PGK1 | phosphoglycerate kinase | -2.5973 | -6.1 | 0.0027821 |
| YLR259C | HSP60 | single-stranded DNA binding | -2.5663 | -5.9 | 0.0284376 |
| YLR134W | PDC5 | pyruvate decarboxylase | -2.5590 | -5.9 | 0.0105047 |
| YOL058W | ARG1 | argininosuccinate synthase | -2.5527 | -5.9 | 0.270517 |
| YGR240C | PFK1 | 6-phosphofructokinase | -2.5485 | -5.9 | 0.0025216 |
| YGR094W | VAS1 | valine-tRNA ligase | -2.5400 | -5.8 | 0.0021247 |
| YHR092C | HXT4 | glucose transporter | -2.5200 | -5.7 | 0.0501148 |
| YGL055W | OLE1 | stearyl-CoA 9-desaturase | -2.5098 | -5.7 | 0.0029415 |
| YGL008C | PMA1 | hydrogen-exporting ATPase , phosphorylative mechanism | -2.4883 | -5.6 | 0.141586 |
| YBR144C | | | -2.4625 | -5.5 | 0.0134941 |
| YGR192C | TDH3 | glyceraldehyde-3-phosphate dehydrogenase | -2.4508 | -5.5 | 0.0056974 |
| YLR044C | PDC1 | pyruvate decarboxylase | -2.4458 | -5.4 | 0.0013079 |
| YHL032C | GUT1 | glycerol kinase | -2.4197 | -5.4 | 0.0316138 |
| YAL037C-B | | | -2.3870 | -5.2 | 0.0004363 |
| YCR013C | | | -2.3687 | -5.2 | 0.0360705 |
| YBR145W | ADH5 | alcohol dehydrogenase | -2.2932 | -4.9 | 0.0057532 |
| YHR094C | HXT1 | glucose transporter | -2.2655 | -4.8 | 0.0067767 |
| YJL052W | TDH1 | glyceraldehyde-3-phosphate dehydrogenase | -2.2213 | -4.7 | 0.0057361 |
| YGR234W | YHB1 | nitric oxide reductase | -2.2198 | -4.7 | 0.010346 |
| YJL005W | CYR1 | adenylate cyclase | -2.2127 | -4.6 | 0.0115928 |
| YDR245W | MNN10 | alpha-1,6-mannosyltransferase | -2.1832 | -4.5 | 0.0208124 |
| YLR056W | ERG3 | C-5 sterol desaturase | -2.1518 | -4.4 | 0.288339 |
| YBL100C | | | -2.1470 | -4.4 | 0.0145585 |
| YIL171W | | unknown | -2.1300 | -4.4 | 0.0214632 |
| YPL028W | ERG10 | acetyl-CoA C-acetyltransferase | -2.1157 | -4.3 | 0.0386839 |
| YDR357C | | unknown | -2.1038 | -4.3 | 0.0132799 |
| YLL066W-A | | | -2.0112 | -4.0 | 0.0062855 |
| YJL045W | | succinate dehydrogenase (ubiquinone) | -2.0098 | -4.0 | 0.0050414 |
| Q0250 | COX2 | cytochrome-c oxidase | -2.0087 | -4.0 | 0.0002256 |
| YEL077C | | helicase | -1.9065 | -3.7 | 0.004981 |
| YBL030C | PET9 | ATP:ADP antiporter | -1.8978 | -3.7 | 0.0021073 |
| YKL148C | SDH1 | succinate dehydrogenase (ubiquinone) | -1.8787 | -3.7 | 0.0072915 |
| YPL187W | MF(ALPHA)1 | mating pheromone | -1.8728 | -3.7 | 0.003322 |
| YGL089C | MF(ALPHA)2 | mating pheromone | -1.8093 | -3.5 | 0.0003334 |
| YIR036C | | unknown | -1.7878 | -3.5 | 0.0096615 |
| YNL055C | POR1 | voltage-gated ion-selective channel | -1.7758 | -3.4 | 0.0041243 |
| YBL099W | ATP1 | hydrogen-transporting ATP synthase , rotational mechanism | -1.7740 | -3.4 | 0.0013101 |
| YAL038W | CDC19 | pyruvate kinase | -1.7672 | -3.4 | 0.0026264 |
| YKL060C | FBA1 | fructose-bisphosphate aldolase | -1.7640 | -3.4 | 0.0057646 |
| YJR048W | CYC1 | electron carrier | -1.7638 | -3.4 | 0.0043225 |
| YFL018C | LPD1 | glycine dehydrogenase (decarboxylating) * | -1.7500 | -3.4 | 0.0069931 |
| YOR187W | TUF1 | GTPase * | -1.7400 | -3.3 | 0.0161058 |
| YBL113W-A | | | -1.7173 | -3.3 | 0.0059674 |
| YDR050C | TPI1 | triose-phosphate isomerase | -1.7088 | -3.3 | 0.001759 |
| YNL135C | FPR1 | peptidyl-prolyl cis-trans isomerase | -1.7083 | -3.3 | 0.0016081 |
| YML110C | COQ5 | 2-hexaprenyl-6-methoxy-1,4-benzoquinone methyltransferase | -1.7035 | -3.3 | 0.0241298 |
| YGR244C | LSC2 | succinate-CoA ligase (ADP-forming) | -1.6993 | -3.2 | 0.0146377 |
| YMR304C-A | | | -1.6962 | -3.2 | 0.0422933 |
| YJR009C | TDH2 | glyceraldehyde-3-phosphate dehydrogenase | -1.6778 | -3.2 | 0.0663943 |
| YJR016C | ILV3 | dihydroxy-acid dehydratase | -1.6523 | -3.1 | 0.0021943 |
| YBR085W | AAC3 | ATP:ADP antiporter | -1.6435 | -3.1 | 0.0005171 |
| YGR287C | | hydrolase , hydrolyzing O-glycosyl compounds | -1.6390 | -3.1 | 0.116654 |
| YPL127C | HHO1 | DNA binding | -1.6355 | -3.1 | 0.0336075 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YDL055C | PSA1 | mannose-1-phosphate guanylyltransferase | -1.6302 | -3.1 | 0.012652 |
| YPL262W | FUM1 | fumarate hydratase | -1.6232 | -3.1 | 0.0035527 |
| YPR204W | | DNA helicase | -1.6188 | -3.1 | 0.0694302 |
| YJR120W | | unknown | -1.6173 | -3.1 | 0.0027704 |
| YKL152C | GPM1 | phosphoglycerate mutase | -1.6137 | -3.1 | 0.045952 |
| YNL339C | YRF1-6 | DNA helicase | -1.5880 | -3.0 | 0.0031108 |
| YNL052W | COX5A | cytochrome-c oxidase | -1.5753 | -3.0 | 0.0040669 |
| YCR074C | | | -1.5665 | -3.0 | 0.005856 |
| TMR307W | | | -1.5593 | -2.9 | 0.0138625 |
| YKL085W | MDH1 | L-malate dehydrogenase | -1.5432 | -2.9 | 0.0049448 |
| YJL214W | HXT8 | glucose transporter | -1.5372 | -2.9 | 0.0406361 |
| YOR133W | EFT1 | translation elongation factor | -1.5342 | -2.9 | 0.0057968 |
| YMR145C | NDE1 | NADH dehydrogenase | -1.5298 | -2.9 | 0.0080462 |
| YGR279C | SCW4 | glucosidase | -1.5277 | -2.9 | 0.0421182 |
| YBL015W | ACH1 | acetyl-CoA hydrolase | -1.5243 | -2.9 | 0.001791 |
| YOR230W | WTM1 | transcription corepressor | -1.5232 | -2.9 | 0.003855 |
| YHR218W | | unknown | -1.5163 | -2.9 | 0.021105 |
| YGL156W | AMS1 | alpha-mannosidase | -1.5097 | -2.8 | 0.0833606 |
| YIR021W | MRS1 | RNA binding | -1.5072 | -2.8 | 0.146316 |
| YGL179C | TOS3 | protein kinase | -1.5000 | -2.8 | 0.0294567 |
| YJR077C | MIR1 | inorganic phosphate transporter | -1.4997 | -2.8 | 0.0078868 |
| YGL202W | ARO8 | aromatic-amino-acid transaminase | -1.4847 | -2.8 | 0.0277173 |
| YNL009W | IDP3 | isocitrate dehydrogenase (NADP+) | -1.4800 | -2.8 | 0.0153217 |
| YBL112C | | unknown | -1.4675 | -2.8 | 0.0238395 |
| YEL030W | ECM10 | unknown | -1.4557 | -2.7 | 0.0147754 |
| YLR420W | URA4 | dihydroorotase | -1.4477 | -2.7 | 0.117124 |
| YNL072W | RNH201 | ribonuclease H | -1.4187 | -2.7 | 0.0114425 |
| YLL026W | HSP104 | chaperone binding | -1.4185 | -2.7 | 0.0789263 |
| YGL253W | HXK2 | hexokinase | -1.4103 | -2.7 | 0.0165731 |
| YNL053W | MSG5 | prenylated protein tyrosine phosphatase | -1.4002 | -2.6 | 0.0045382 |
| YMR056C | AAC1 | ATP:ADP antiporter | -1.3943 | -2.6 | 0.0132202 |
| YGR143W | SKN1 | glucosidase | -1.3917 | -2.6 | 0.0102354 |
| YHR051W | COX6 | cytochrome-c oxidase | -1.3872 | -2.6 | 0.002703 |
| YML028W | TSA1 | thioredoxin peroxidase | -1.3855 | -2.6 | 0.0066019 |
| YDL021W | GPM2 | unknown | -1.3822 | -2.6 | 0.0126967 |
| YJR103W | URA8 | CTP synthase | -1.3815 | -2.6 | 0.0375282 |
| YEL066W | HPA3 | histone acetyltransferase | -1.3765 | -2.6 | 0.0015326 |
| YLR174W | IDP2 | isocitrate dehydrogenase (NADP+) | -1.3750 | -2.6 | 0.0037159 |
| YJL217W | | unknown | -1.3702 | -2.6 | 0.0006493 |
| YHL034C | SBP1 | RNA binding | -1.3697 | -2.6 | 0.0234807 |
| YEL024W | RIP1 | ubiquinol-cytochrome-c reductase | -1.3682 | -2.6 | 0.0128749 |
| YNR036C | | structural constituent of ribosome | -1.3630 | -2.6 | 0.0271336 |
| YMR199W | CLN1 | cyclin-dependent protein kinase regulator | -1.3627 | -2.6 | 0.0350652 |
| YJR148W | BAT2 | branched-chain-amino-acid transaminase | -1.3610 | -2.6 | 0.0177124 |
| YGL191W | COX13 | enzyme regulator * | -1.3572 | -2.6 | 0.0054378 |
| YHL050C | | helicase | -1.3523 | -2.6 | 0.0358242 |
| YOR393W | ERR1 | phosphopyruvate hydratase | -1.3355 | -2.5 | 0.220241 |
| YOR142W | LSC1 | succinate-CoA ligase (ADP-forming) | -1.3352 | -2.5 | 0.0005867 |
| YMR015C | ERG5 | C-22 sterol desaturase | -1.3238 | -2.5 | 0.0469431 |
| YBR026C | ETR1 | enoyl-[acyl-carrier protein] reductase | -1.3205 | -2.5 | 7.395E-06 |
| YGR191W | HIP1 | L-histidine transporter | -1.3102 | -2.5 | 0.0027607 |
| YDR385W | EFT2 | translation elongation factor | -1.3068 | -2.5 | 0.0039284 |
| YJR031C | GEA1 | ARF guanyl-nucleotide exchange factor | -1.2995 | -2.5 | 0.0164891 |
| YFR053C | HXK1 | hexokinase | -1.2933 | -2.5 | 0.0070769 |
| YHR219W | | unknown | -1.2927 | -2.4 | 0.0065523 |
| YGL062W | PYC1 | pyruvate carboxylase | -1.2910 | -2.4 | 0.019084 |
| YLR395C | COX8 | cytochrome-c oxidase | -1.2893 | -2.4 | 0.0146455 |
| YDR202C | RAV2 | unknown | -1.2888 | -2.4 | 0.0007558 |
| YDR119W | | unknown | -1.2875 | -2.4 | 0.0012483 |
| YLL041C | SDH2 | succinate dehydrogenase (ubiquinone) | -1.2683 | -2.4 | 0.0082625 |
| YDL185W | TFP1 | hydrogen-transporting ATPase, rotational mechanism* | -1.2658 | -2.4 | 0.0237687 |
| YER023W | PRO3 | pyrroline-5-carboxylate reductase | -1.2652 | -2.4 | 0.0012985 |
| YNL220W | ADE12 | adenylosuccinate synthase | -1.2637 | -2.4 | 0.0001225 |
| YGL105W | ARC1 | tRNA binding | -1.2623 | -2.4 | 0.0085279 |
| YKL097W-A | | | -1.2552 | -2.4 | 0.0162569 |
| YKL067W | YNK1 | nucleoside diphosphate kinase | -1.2533 | -2.4 | 0.0032047 |
| YKL153W | | | -1.2488 | -2.4 | 0.0168587 |
| YER178W | PDA1 | pyruvate dehydrogenase (acetyl-transferring) | -1.2483 | -2.4 | 0.0091187 |
| YPL134C | ODC1 | intracellular transporter | -1.2475 | -2.4 | 0.0061971 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YKL217W | JEN1 | lactate transporter | -1.2455 | -2.4 | 0.0558608 |
| YEL039C | CYC7 | electron carrier | -1.2420 | -2.4 | 0.0019586 |
| YHR212C | | | -1.2417 | -2.4 | 0.145285 |
| YCL025C | AGP1 | amino acid transporter | -1.2355 | -2.4 | 0.0037559 |
| YGL188C | | | -1.2317 | -2.3 | 0.0336314 |
| YFL039C | ACT1 | structural constituent of cytoskeleton | -1.2285 | -2.3 | 0.0084709 |
| YDR453C | TSA2 | thioredoxin peroxidase | -1.2272 | -2.3 | 0.0077252 |
| YPL078C | ATP4 | structural molecule * | -1.2218 | -2.3 | 0.0033471 |
| YBL098W | BNA4 | kynurenine 3-monooxygenase | -1.2158 | -2.3 | 0.0178262 |
| YNR001C | CIT1 | citrate (Si)-synthase | -1.2085 | -2.3 | 0.0021957 |
| YBR078W | ECM33 | unknown | -1.2015 | -2.3 | 0.0015443 |
| YMR190C | SGS1 | ATP-dependent DNA helicase | -1.1968 | -2.3 | 0.0018184 |
| YMR208W | ERG12 | mevalonate kinase | -1.1950 | -2.3 | 0.0526759 |
| YDR273W | DON1 | unknown | -1.1930 | -2.3 | 0.0052203 |
| YER053C | PIC2 | inorganic phosphate transporter | -1.1862 | -2.3 | 0.002229 |
| YDL100C | GET3 | ATPase | -1.1862 | -2.3 | 0.0198745 |
| YIR037W | HYR1 | thioredoxin peroxidase * | -1.1848 | -2.3 | 0.0324715 |
| YNL103W | MET4 | transcription coactivator | -1.1798 | -2.3 | 0.0017762 |
| YIR035C | | unknown | -1.1687 | -2.2 | 0.013592 |
| YMR297W | PRC1 | carboxypeptidase C | -1.1672 | -2.2 | 0.0058152 |
| YER133W | GLC7 | protein phosphatase type 1 | -1.1610 | -2.2 | 0.0084142 |
| YGR292W | MAL12 | alpha-glucosidase | -1.1568 | -2.2 | 0.0960076 |
| YJR139C | HOM6 | homoserine dehydrogenase | -1.1497 | -2.2 | 0.0071506 |
| YHR007C | ERG11 | sterol 14-demethylase | -1.1395 | -2.2 | 0.0049116 |
| YOR084W | | lipase | -1.1385 | -2.2 | 0.0039637 |
| YDL126C | CDC48 | ATPase | -1.1323 | -2.2 | 0.0240621 |
| YDR148C | KGD2 | dihydrolipoyllysine-residue succinyltransferase | -1.1323 | -2.2 | 0.0018843 |
| YDR284C | DPP1 | phosphatidate phosphatase * | -1.1293 | -2.2 | 0.0041214 |
| YKL176C | LST4 | protein transporter | -1.1257 | -2.2 | 0.339576 |
| YBR244W | GPX2 | glutathione peroxidase * | -1.1227 | -2.2 | 0.0040126 |
| YKL141W | SDH3 | succinate dehydrogenase (ubiquinone) | -1.1197 | -2.2 | 0.0064126 |
| YLR460C | | unknown | -1.1190 | -2.2 | 0.0404649 |
| YDL215C | GDH2 | glutamate dehydrogenase | -1.1168 | -2.2 | 0.0244043 |
| YBL001C | ECM15 | unknown | -1.1123 | -2.2 | 0.0186059 |
| YMR215W | GAS3 | 1,3-beta-glucanosyltransferase | -1.1123 | -2.2 | 0.0490475 |
| YKL066W | | | -1.1013 | -2.1 | 0.0051052 |
| YML132W | COS3 | protein binding | -1.0953 | -2.1 | 0.0048383 |
| YKL192C | ACP1 | acyl carrier | -1.0917 | -2.1 | 0.014706 |
| YLL018C | DPS1 | RNA binding | -1.0917 | -2.1 | 0.0193234 |
| YKR046C | PET10 | unknown | -1.0903 | -2.1 | 0.0395881 |
| YDL210W | UGA4 | putrescine transporter | -1.0775 | -2.1 | 0.083624 |
| YBR185C | MBA1 | unknown | -1.0762 | -2.1 | 0.0167391 |
| YBL075C | SSA3 | ATPase * | -1.0710 | -2.1 | 0.0055408 |
| YER152C | | unknown | -1.0665 | -2.1 | 0.115109 |
| YKL157W | APE2 | leucyl aminopeptidase | -1.0643 | -2.1 | 0.000473 |
| YHR008C | SOD2 | manganese superoxide dismutase | -1.0640 | -2.1 | 0.0058543 |
| YGL200C | EMP24 | unknown | -1.0520 | -2.1 | 0.0414524 |
| YKL174C | TPO5 | polyamine transporter | -1.0508 | -2.1 | 0.128647 |
| YML078W | CPR3 | peptidyl-prolyl cis-trans isomerase | -1.0508 | -2.1 | 0.0186655 |
| YKL069W | | unknown | -1.0442 | -2.1 | 0.112305 |
| YEL009C | GCN4 | DNA binding | -1.0337 | -2.0 | 0.0097289 |
| YDR438W | | unknown | -1.0330 | -2.0 | 0.002288 |
| YDL067C | COX9 | cytochrome-c oxidase | -1.0290 | -2.0 | 0.0015579 |
| YDR200C | VPS64 | unknown | -1.0273 | -2.0 | 0.242927 |
| YBR067C | TIP1 | structural constituent of cell wall* | -1.0262 | -2.0 | 0.0311461 |
| YHL034W-A | | | -1.0245 | -2.0 | 0.0514341 |
| YHR126C | | unknown | -1.0212 | -2.0 | 0.0043803 |
| YPL082C | MOT1 | ATPase | -1.0182 | -2.0 | 0.0625089 |
| YCRX18C | | | -1.0172 | -2.0 | 0.0930517 |
| YFL062W | COS4 | unknown | -1.0137 | -2.0 | 0.0043981 |
| YHR214W | | unknown | -1.0113 | -2.0 | 0.0006442 |
| YER011W | TIR1 | structural constituent of cell wall | -1.0028 | -2.0 | 0.0148515 |
| YHR028C | DAP2 | dipeptidyl-peptidase and tripeptidyl-peptidase | -1.0017 | -2.0 | 0.0039903 |
| YKL160W | ELF1 | RNA polymerase II transcription elongation factor | -0.9982 | -2.0 | 0.0409732 |
| YGR167W | CLC1 | structural molecule | -0.9933 | -2.0 | 0.0137311 |
| YLR304C | ACO1 | aconitate hydratase | -0.9883 | -2.0 | 0.0341339 |
| YPL080C | | | -0.9882 | -2.0 | 0.0301871 |
| YBL045C | COR1 | ubiquinol-cytochrome-c reductase | -0.9868 | -2.0 | 0.0011312 |
| YHR001W-A | QCR10 | ubiquinol-cytochrome-c reductase | -0.9818 | -2.0 | 0.0243084 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YBL102W | SFT2 | unknown | -0.9788 | -2.0 | 0.0469287 |
| YJL068C | | S-formylglutathione hydrolase | -0.9782 | -2.0 | 0.0029145 |
| YBL078C | ATG8 | microtubule binding | -0.9772 | -2.0 | 0.0238878 |
| YIL166C | | transporter | -0.9752 | -2.0 | 0.0027811 |
| YGR175C | ERG1 | squalene monooxygenase * | -0.9740 | -2.0 | 0.0041998 |
| YAR018C | KIN3 | protein kinase | -0.9685 | -2.0 | 0.0155214 |
| YLR150W | STM1 | DNA binding | -0.9665 | -2.0 | 0.0482563 |
| YIL125W | KGD1 | oxoglutarate dehydrogenase (succinyl-transferring) | -0.9652 | -2.0 | 0.0669744 |
| YFL049W | SWP82 | transcriptional activator | -0.9628 | -1.9 | 0.0244383 |
| YDL066W | IDP1 | isocitrate dehydrogenase (NADP+) | -0.9600 | -1.9 | 0.0016617 |
| YIL057C | | unknown | -0.9572 | -1.9 | 0.023131 |
| YKR013W | PRY2 | unknown | -0.9455 | -1.9 | 0.247454 |
| YGR121C | MEP1 | ammonium transporter | -0.9423 | -1.9 | 0.368745 |
| YMR205C | PFK2 | 6-phosphofructokinase | -0.9413 | -1.9 | 0.0038085 |
| YLR303W | MET17 | cysteine synthase * | -0.9403 | -1.9 | 0.0102335 |
| YDR327W | | | -0.9383 | -1.9 | 0.0063162 |
| YPR020W | ATP20 | structural molecule | -0.9380 | -1.9 | 0.0218309 |
| YNL071W | LAT1 | dihydrolipoyllysine-residue acetyltransferase | -0.9350 | -1.9 | 0.01109 |
| YJL066C | MPM1 | unknown | -0.9325 | -1.9 | 0.0049339 |
| YMR206W | | unknown | -0.9315 | -1.9 | 0.0173931 |
| YNL336W | COS1 | unknown | -0.9305 | -1.9 | 0.0080794 |
| YDR047W | HEM12 | uroporphyrinogen decarboxylase | -0.9270 | -1.9 | 0.0240862 |
| YER003C | PMI40 | mannose-6-phosphate isomerase | -0.9263 | -1.9 | 0.0071212 |
| YFL038C | YPT1 | GTPase | -0.9255 | -1.9 | 0.0057113 |
| YIL157C | FMP35 | unknown | -0.9255 | -1.9 | 0.0011721 |
| YBR025C | | unknown | -0.9253 | -1.9 | 0.0007142 |
| YBR243C | ALG7 | UDP-N-acetylglucosamine-dolichyl-phosphate N-acetylglucosyl transferase | -0.9253 | -1.9 | 0.0358139 |
| YLL052C | AQY2 | water channel | -0.9215 | -1.9 | 0.037856 |
| YBR148W | YSW1 | unknown | -0.9202 | -1.9 | 0.0274711 |
| YCL057W | PRD1 | metalloendopeptidase * | -0.9110 | -1.9 | 0.0505244 |
| YGR246C | BRF1 | RNA polymerase III transcription factor | -0.9107 | -1.9 | 0.0120764 |
| YDR193W | | | -0.9100 | -1.9 | 0.0027404 |
| YDL120W | YFH1 | enzyme regulator | -0.9063 | -1.9 | 0.0008325 |
| YDL078C | MDH3 | L-malate dehydrogenase | -0.9043 | -1.9 | 0.0007158 |
| YAL062W | GDH3 | glutamate dehydrogenase | -0.9028 | -1.9 | 0.0829544 |
| YGR295C | COS6 | unknown | -0.8932 | -1.9 | 0.0233409 |
| YGR142W | BTN2 | unknown | -0.8875 | -1.8 | 0.127621 |
| YJL213W | | unknown | -0.8865 | -1.8 | 0.0070708 |
| YDR032C | PST2 | unknown | -0.8860 | -1.8 | 0.0097202 |
| YEL065W | SIT1 | siderophore-iron (ferrioxamine) uptake transporter | -0.8780 | -1.8 | 0.0025091 |
| YJR080C | FMP26 | unknown | -0.8747 | -1.8 | 0.0051612 |
| YDR007W | TRP1 | phosphoribosylanthranilate isomerase | -0.8745 | -1.8 | 0.0511363 |
| YJL048C | UBX6 | unknown | -0.8720 | -1.8 | 0.0006909 |
| YLR231C | BNA5 | kynureninase | -0.8720 | -1.8 | 0.135077 |
| YIL123W | SIM1 | unknown | -0.8695 | -1.8 | 0.0469092 |
| YOR007C | SGT2 | unknown | -0.8687 | -1.8 | 0.0062891 |
| YHR057C | CPR2 | peptidyl-prolyl cis-trans isomerase | -0.8682 | -1.8 | 0.178201 |
| YKL216W | URA1 | dihydroorotate dehydrogenase | -0.8667 | -1.8 | 0.0028603 |
| YHR018C | ARG4 | argininosuccinate lyase | -0.8643 | -1.8 | 0.0131009 |
| YAL008W | FUN14 | unknown | -0.8628 | -1.8 | 0.0025063 |
| YKR020W | VPS51 | protein binding | -0.8627 | -1.8 | 0.175689 |
| YGR132C | PHB1 | unknown | -0.8623 | -1.8 | 0.0238722 |
| YIR044C | | unknown | -0.8613 | -1.8 | 0.0067745 |
| YML039W | | | -0.8602 | -1.8 | 0.113936 |
| YLR294C | | | -0.8593 | -1.8 | 0.0185024 |
| YFL005W | SEC4 | GTPase | -0.8590 | -1.8 | 0.0094263 |
| YGL026C | TRP5 | tryptophan synthase | -0.8500 | -1.8 | 0.11978 |
| YFR044C | | unknown | -0.8477 | -1.8 | 0.0633565 |
| YBR294W | SUL1 | sulfate transporter | -0.8468 | -1.8 | 0.0223382 |
| YGR021W | | unknown | -0.8463 | -1.8 | 0.0452241 |
| YML120C | NDI1 | NADH dehydrogenase (ubiquinone) | -0.8417 | -1.8 | 0.0094716 |
| YDR268W | MSW1 | tryptophan-tRNA ligase | -0.8390 | -1.8 | 0.008626 |
| YGL164C | YRB30 | protein binding | -0.8355 | -1.8 | 0.106843 |
| YMR074C | | unknown | -0.8355 | -1.8 | 0.0106242 |
| YDL181W | INH1 | enzyme inhibitor | -0.8330 | -1.8 | 0.0336067 |
| YGL068W | MNP1 | unknown | -0.8325 | -1.8 | 0.0212398 |
| YNL037C | IDH1 | isocitrate dehydrogenase (NAD+) | -0.8230 | -1.8 | 0.0188632 |
| YJR073C | OPI3 | phosphatidyl-N-methylethanolamine N-methyltransferase | -0.8185 | -1.8 | 0.0331786 |
| YGR253C | PUP2 | endopeptidase | -0.8150 | -1.8 | 0.000143 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YBR039W | ATP3 | hydrogen-transporting ATP synthase , rotational mechanism | -0.8148 | -1.8 | 0.0102082 |
| YJR109C | CPA2 | carbamoyl-phosphate synthase (glutamine-hydrolyzing) | -0.8132 | -1.8 | 0.0905807 |
| YER010C | | unknown | -0.8112 | -1.8 | 0.444645 |
| YPR033C | HTS1 | histidine-tRNA ligase | -0.8100 | -1.8 | 0.0051138 |
| YNL144C | | unknown | -0.8055 | -1.7 | 0.183034 |
| YEL052W | AFG1 | ATPase | -0.8032 | -1.7 | 0.0369059 |
| YDL237W | | unknown | -0.8028 | -1.7 | 0.0107521 |
| YJL079C | PRY1 | unknown | -0.7997 | -1.7 | 0.0032116 |
| YKR076W | ECM4 | unknown | -0.7987 | -1.7 | 0.0189093 |
| YGL148W | ARO2 | chorismate synthase * | -0.7973 | -1.7 | 0.062767 |
| YCR021C | HSP30 | unknown | -0.7970 | -1.7 | 0.0112727 |
| YFR003C | YPI1 | protein phosphatase inhibitor | -0.7953 | -1.7 | 0.0347147 |
| YGR183C | QCR9 | ubiquinol-cytochrome-c reductase | -0.7947 | -1.7 | 0.029432 |
| YGL004C | RPN14 | unknown | -0.7943 | -1.7 | 0.0302875 |
| YDL248W | COS7 | receptor | -0.7927 | -1.7 | 0.0090428 |
| YMR108W | ILV2 | acetolactate synthase * | -0.7923 | -1.7 | 0.0066842 |
| YNL057W | | | -0.7885 | -1.7 | 0.139776 |
| YDR163W | CWC15 | unknown | -0.7867 | -1.7 | 0.0265685 |
| YBR299W | MAL32 | alpha-glucosidase | -0.7815 | -1.7 | 0.025647 |
| YGR182C | | | -0.7792 | -1.7 | 0.0436314 |
| YDL195W | SEC31 | structural molecule | -0.7775 | -1.7 | 0.0054941 |
| YPR088C | SRP54 | signal sequence binding | -0.7770 | -1.7 | 0.0858451 |
| YHR074W | QNS1 | hydrolase , acting on carbon-nitrogen (but not peptide) bonds, i | -0.7753 | -1.7 | 0.378315 |
| YHL048W | COS8 | unknown | -0.7737 | -1.7 | 0.0130009 |
| YMR276W | DSK2 | protein binding | -0.7735 | -1.7 | 0.0480635 |
| YAL055W | PEX22 | unknown | -0.7727 | -1.7 | 0.0669845 |
| YIL136W | OM45 | unknown | -0.7725 | -1.7 | 0.0169707 |
| YHL019W-A | | | -0.7703 | -1.7 | 0.18253 |
| YJL001W | PRE3 | endopeptidase | -0.7703 | -1.7 | 0.0377976 |
| YDR031W | | unknown | -0.7692 | -1.7 | 0.0012387 |
| YPL265W | DIP5 | amino acid transporter | -0.7660 | -1.7 | 0.0286061 |
| YJL207C | LAA1 | unknown | -0.7643 | -1.7 | 0.0315624 |
| YIL124W | AYR1 | acylglycerone-phosphate reductase | -0.7625 | -1.7 | 0.0880242 |
| YDL174C | DLD1 | D-lactate dehydrogenase (cytochrome) | -0.7615 | -1.7 | 0.0139759 |
| YKR048C | NAP1 | protein binding | -0.7610 | -1.7 | 0.0386136 |
| YIR043C | | unknown | -0.7595 | -1.7 | 0.0033023 |
| YBR221C | PDB1 | pyruvate dehydrogenase (acetyl-transferring) | -0.7580 | -1.7 | 0.0147966 |
| YER056C | FCY2 | cytosine-purine permease * | -0.7563 | -1.7 | 0.108878 |
| YCL058C | | | -0.7530 | -1.7 | 0.0382909 |
| YIR038C | GTT1 | glutathione transferase | -0.7527 | -1.7 | 0.0203948 |
| YBR219C | | unknown | -0.7518 | -1.7 | 0.0233464 |
| YKL164C | PIR1 | structural constituent of cell wall | -0.7518 | -1.7 | 0.0092288 |
| YCL009C | ILV6 | enzyme regulator * | -0.7505 | -1.7 | 0.0087365 |
| YDL171C | GLT1 | glutamate synthase (NADH) | -0.7493 | -1.7 | 0.0157805 |
| YPL048W | CAM1 | translation elongation factor | -0.7467 | -1.7 | 0.0047668 |
| YNR018W | | unknown | -0.7445 | -1.7 | 0.0393113 |
| YPR156C | TPO3 | spermine transporter | -0.7422 | -1.7 | 0.058331 |
| YDR157W | | | -0.7415 | -1.7 | 0.0330321 |
| YOL071W | EMI5 | unknown | -0.7372 | -1.7 | 0.0220893 |
| YDR388W | RVS167 | cytoskeletal protein binding | -0.7368 | -1.7 | 0.0366519 |
| YNL064C | YDJ1 | chaperone regulator * | -0.7345 | -1.7 | 0.0410423 |
| YBR094W | | unknown | -0.7335 | -1.7 | 0.001266 |
| YCL030C | HIS4 | phosphoribosyl-ATP diphosphatase | -0.7307 | -1.7 | 0.0471703 |
| YGL048C | RPT6 | ATPase * | -0.7280 | -1.7 | 0.0341018 |
| YLR028C | ADE16 | IMP cyclohydrolase | -0.7262 | -1.7 | 0.0107976 |
| YIL137C | TMA108 | metalloendopeptidase | -0.7255 | -1.7 | 0.125156 |
| YCR079W | | phosphoprotein phosphatase | -0.7242 | -1.7 | 0.0464033 |
| YDL192W | ARF1 | GTPase | -0.7237 | -1.7 | 0.0199873 |
| YOR243C | PUS7 | pseudouridine synthase | -0.7217 | -1.6 | 0.0269627 |
| YOR215C | | unknown | -0.7203 | -1.6 | 0.0006178 |
| YDR129C | SAC6 | protein binding | -0.7160 | -1.6 | 0.0715468 |
| YCR058C | | | -0.7150 | -1.6 | 0.266776 |
| YER004W | FMP52 | unknown | -0.7143 | -1.6 | 0.0035528 |
| YFR031C-A | RPL2A | structural constituent of ribosome | -0.7118 | -1.6 | 0.0094638 |
| YHR037W | PUT2 | l-pyrroline-5-carboxylate dehydrogenase | -0.7110 | -1.6 | 0.0044512 |
| YKL103C | LAP4 | aminopeptidase I | -0.7053 | -1.6 | 0.0286373 |
| YER103W | SSA4 | ATPase * | -0.7045 | -1.6 | 0.0973909 |
| YCR004C | YCP4 | electron carrier | -0.7020 | -1.6 | 0.0363207 |
| YCR014C | POL4 | beta DNA polymerase | -0.7017 | -1.6 | 0.140327 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YBR046C | ZTA1 | unknown | -0.6998 | -1.6 | 7.841E-06 |
| YLR248W | RCK2 | protein serine/threonine kinase | -0.6997 | -1.6 | 0.0040415 |
| YHR214W-A | | | -0.6997 | -1.6 | 0.080264 |
| YGR231C | PHB2 | unknown | -0.6987 | -1.6 | 0.0250179 |
| YPL270W | MDL2 | ATPase , coupled to transmembrane movement of substances | -0.6985 | -1.6 | 0.0241096 |
| YDR310C | SUM1 | transcriptional repressor | -0.6975 | -1.6 | 0.0007879 |
| YDL137W | ARF2 | GTPase | -0.6948 | -1.6 | 0.0250585 |
| YFR024C | | | -0.6937 | -1.6 | 0.0249951 |
| YOR289W | | unknown | -0.6935 | -1.6 | 0.051487 |
| YGL115W | SNF4 | protein kinase activator | -0.6917 | -1.6 | 0.0393142 |
| YLR257W | | unknown | -0.6860 | -1.6 | 0.0098596 |
| YEL025C | | unknown | -0.6858 | -1.6 | 0.347913 |
| YJR001W | AVT1 | neutral amino acid transporter | -0.6825 | -1.6 | 0.0168642 |
| YPL031C | PHO85 | cyclin-dependent protein kinase | -0.6825 | -1.6 | 0.0059765 |
| YJL081C | ARP4 | chromatin binding | -0.6797 | -1.6 | 0.0894538 |
| YDR325W | YCG1 | unknown | -0.6795 | -1.6 | 0.0257903 |
| YPL274W | SAM3 | S-adenosylmethionine transporter | -0.6787 | -1.6 | 0.0428522 |
| YBR298C | MAL31 | alpha-glucoside:hydrogen symporter | -0.6783 | -1.6 | 0.0030428 |
| YGL121C | GPG1 | signal transducer | -0.6750 | -1.6 | 0.0086372 |
| YLL024C | SSA2 | ATPase * | -0.6732 | -1.6 | 0.023702 |
| YIL065C | FIS1 | unknown | -0.6730 | -1.6 | 0.0232047 |
| YEL032W | MCM3 | chromatin binding | -0.6725 | -1.6 | 0.0296826 |
| YGR038W | ORM1 | unknown | -0.6725 | -1.6 | 0.0568904 |
| YJL034W | KAR2 | ATPase * | -0.6698 | -1.6 | 0.0194184 |
| YER012W | PRE1 | endopeptidase | -0.6693 | -1.6 | 0.0629897 |
| YPL034W | | unknown | -0.6687 | -1.6 | 0.373552 |
| YFL045C | SEC53 | phosphomannomutase | -0.6667 | -1.6 | 0.0270839 |
| YGR250C | | RNA binding | -0.6667 | -1.6 | 0.000825 |
| YBR139W | | serine hydrolase * | -0.6622 | -1.6 | 0.0049168 |
| YJR023C | | | -0.6613 | -1.6 | 0.0192836 |
| YCL018W | LEU2 | 3-isopropylmalate dehydrogenase | -0.6597 | -1.6 | 0.0063544 |
| YIL078W | THS1 | threonine-tRNA ligase | -0.6595 | -1.6 | 0.0125413 |
| YAL023C | PMT2 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.6578 | -1.6 | 0.0073141 |
| YKL200C | | | -0.6575 | -1.6 | 0.0263667 |
| YPR024W | YME1 | ATP-dependent peptidase | -0.6548 | -1.6 | 0.191781 |
| YLR354C | TAL1 | transaldolase | -0.6543 | -1.6 | 0.0634906 |
| YML106W | URA5 | orotate phosphoribosyltransferase | -0.6480 | -1.6 | 0.0026807 |
| YPR083W | MDM36 | unknown | -0.6452 | -1.6 | 0.129119 |
| YJR079W | | unknown | -0.6433 | -1.6 | 0.13572 |
| YLR095C | IOC2 | protein binding | -0.6368 | -1.6 | 0.0097903 |
| YDL029W | ARP2 | structural constituent of cytoskeleton* | -0.6367 | -1.6 | 0.0048151 |
| YHR117W | TOM71 | protein transporter | -0.6330 | -1.6 | 0.0315424 |
| YPL059W | GRX5 | thiol-disulfide exchange intermediate | -0.6285 | -1.5 | 0.203837 |
| YDR023W | SES1 | serine-tRNA ligase | -0.6282 | -1.5 | 0.007585 |
| YER031C | YPT31 | GTPase | -0.6250 | -1.5 | 0.0240893 |
| YBL003C | HTA2 | DNA binding | -0.6233 | -1.5 | 0.0513053 |
| YBR088C | POL30 | DNA polymerase processivity factor | -0.6232 | -1.5 | 0.0115579 |
| YDR177W | UBC1 | ubiquitin conjugating enzyme | -0.6227 | -1.5 | 0.001769 |
| YML054C | CYB2 | L-lactate dehydrogenase (cytochrome) | -0.6225 | -1.5 | 0.0152554 |
| YER136W | GDI1 | Rab GDP-dissociation inhibitor | -0.6218 | -1.5 | 0.0543204 |
| YIL051C | MMF1 | unknown | -0.6212 | -1.5 | 0.0220292 |
| YFL066C | | unknown | -0.6177 | -1.5 | 0.0117956 |
| YPL282C | PAU22 | unknown | -0.6138 | -1.5 | 0.0428703 |
| YIR022W | SEC11 | signal peptidase | -0.6132 | -1.5 | 0.001555 |
| YGR214W | RPS0A | structural constituent of ribosome | -0.6070 | -1.5 | 0.0203501 |
| YNL100W | | unknown | -0.6058 | -1.5 | 0.0295146 |
| YOL111C | MDY2 | unknown | -0.6058 | -1.5 | 0.0614107 |
| YBR074W | | metalloendopeptidase | -0.6047 | -1.5 | 0.0153701 |
| YMR099C | | unknown | -0.5987 | -1.5 | 0.0688834 |
| YCR011C | ADP1 | ATPase | -0.5898 | -1.5 | 0.0220968 |
| YER091C | MET6 | 5-methyltetrahydropteroyltrimethylglutamate-homocysteine S-methylt | -0.5895 | -1.5 | 0.104279 |
| YER095W | RAD51 | recombinase | -0.5878 | -1.5 | 0.158913 |
| YEL034C-A | | | -0.5863 | -1.5 | 0.0616625 |
| YJL014W | CCT3 | unfolded protein binding | -0.5853 | -1.5 | 0.0123731 |
| YJL012C | VTC4 | unknown | -0.5852 | -1.5 | 0.113078 |
| YBR280C | | unknown | -0.5807 | -1.5 | 0.0332498 |
| YPR035W | GLN1 | glutamate-ammonia ligase | -0.5800 | -1.5 | 0.0130086 |
| YDR529C | QCR7 | ubiquinol-cytochrome-c reductase | -0.5793 | -1.5 | 0.0832114 |
| YOR335C | ALA1 | alanine-tRNA ligase | -0.5790 | -1.5 | 0.145233 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YDR429C | TIF35 | translation initiation factor | -0.5782 | -1.5 | 0.0236947 |
| YGL066W | SGF73 | structural molecule * | -0.5778 | -1.5 | 0.0289986 |
| YCL061C | MRC1 | unknown | -0.5730 | -1.5 | 0.0227256 |
| YMR203W | TOM40 | protein transporter | -0.5712 | -1.5 | 0.0408657 |
| YJL143W | TIM17 | protein transporter | -0.5705 | -1.5 | 0.000366 |
| YDR198C | RKM2 | unknown | -0.5702 | -1.5 | 0.0394924 |
| YLR326W | | unknown | -0.5697 | -1.5 | 0.0382259 |
| YPR028W | YOP1 | protein binding | -0.5677 | -1.5 | 0.116211 |
| YDR496C | PUF6 | specific transcriptional repressor | -0.5652 | -1.5 | 0.0444239 |
| YBR206W | | | -0.5643 | -1.5 | 0.0509919 |
| YDL093W | PMT5 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.5630 | -1.5 | 0.148798 |
| YOR168W | GLN4 | glutamine-tRNA ligase | -0.5627 | -1.5 | 0.0028085 |
| YOR259C | RPT4 | ATPase * | -0.5623 | -1.5 | 0.0217712 |
| YEL042W | GDA1 | guanosine-diphosphatase | -0.5613 | -1.5 | 0.0444184 |
| YJR014W | TMA22 | RNA binding | -0.5605 | -1.5 | 0.0890437 |
| YPR080W | TEF1 | translation elongation factor | -0.5558 | -1.5 | 0.0850207 |
| YKR070W | | unknown | -0.5547 | -1.5 | 0.0165033 |
| YBR121C | GRS1 | glycine-tRNA ligase | -0.5535 | -1.5 | 0.0040589 |
| YEL011W | GLC3 | 1,4-alpha-glucan branching enzyme | -0.5535 | -1.5 | 0.0515006 |
| YCL064C | CHA1 | L-serine ammonia-lyase * | -0.5522 | -1.5 | 0.0077958 |
| YGR271W | SLH1 | RNA helicase | -0.5475 | -1.5 | 0.0124169 |
| YER009W | NTF2 | Ran GTPase binding | -0.5455 | -1.5 | 0.0210792 |
| YOR020C | HSP10 | unfolded protein binding | -0.5433 | -1.5 | 0.03817 |
| YDL007W | RPT2 | ATPase * | -0.5405 | -1.5 | 0.010477 |
| YDR253C | MET32 | DNA binding | -0.5402 | -1.5 | 0.146406 |
| YML012W | ERV25 | unknown | -0.5395 | -1.5 | 0.012581 |
| YEL049W | PAU2 | unknown | -0.5368 | -1.5 | 0.0479029 |
| YJL062W | LAS21 | transferase | -0.5358 | -1.4 | 0.0018635 |
| YLR378C | SEC61 | protein transporter | -0.5355 | -1.4 | 0.052101 |
| YML010W | SPT5 | RNA polymerase II transcription elongation factor | -0.5353 | -1.4 | 0.0234722 |
| YPL181W | CTI6 | transcription factor binding | -0.5332 | -1.4 | 0.0626876 |
| YBR054W | YRO2 | unknown | -0.5330 | -1.4 | 0.0420432 |
| YNL134C | | alcohol dehydrogenase (NADP+) | -0.5302 | -1.4 | 0.0194446 |
| YEL034W | HYP2 | protein binding | -0.5297 | -1.4 | 0.0165189 |
| YFL022C | FRS2 | phenylalanine-tRNA ligase | -0.5258 | -1.4 | 0.0023064 |
| YIL075C | RPN2 | endopeptidase * | -0.5253 | -1.4 | 0.0473779 |
| YFL037W | TUB2 | structural constituent of cytoskeleton | -0.5240 | -1.4 | 0.0165442 |
| YKL109W | HAP4 | transcriptional activator | -0.5240 | -1.4 | 0.0089356 |
| YBR208C | DUR1,2 | allophanate hydrolase * | -0.5230 | -1.4 | 0.0170561 |
| YIL039W | | unknown | -0.5230 | -1.4 | 0.0190342 |
| YDR086C | SSS1 | protein transporter | -0.5228 | -1.4 | 0.0070185 |
| YPR026W | ATH1 | alpha, alpha-trehalase | -0.5203 | -1.4 | 0.0173272 |
| YBR301W | DAN3 | unknown | -0.5182 | -1.4 | 0.0297179 |
| YKL163W | PIR3 | structural constituent of cell wall | -0.5173 | -1.4 | 0.0252202 |
| YJL080C | SCP160 | RNA binding | -0.5172 | -1.4 | 0.0491651 |
| YKR094C | RPL40B | structural constituent of ribosome* | -0.5167 | -1.4 | 0.0231704 |
| YLR249W | YEF3 | translation elongation factor | -0.5167 | -1.4 | 0.078223 |
| YOR394W | PAU21 | unknown | -0.5167 | -1.4 | 0.0543114 |
| YMR104C | YPK2 | protein kinase | -0.5165 | -1.4 | 0.127505 |
| YOR136W | IDH2 | isocitrate dehydrogenase (NAD+) | -0.5142 | -1.4 | 0.0216685 |
| YPL224C | MMT2 | unknown | -0.5138 | -1.4 | 0.0440024 |
| YDL206W | | unknown | -0.5130 | -1.4 | 0.0046788 |
| YOR135C | | | -0.5120 | -1.4 | 0.0647748 |
| YPL179W | PPQ1 | protein serine/threonine phosphatase | -0.5103 | -1.4 | 0.0072955 |
| YKL212W | SAC1 | inositol or phosphatidylinositol phosphatase | -0.5092 | -1.4 | 0.0424713 |
| YNL173C | MDG1 | unknown | -0.5087 | -1.4 | 0.0809204 |
| YDR377W | ATP17 | hydrogen-transporting ATP synthase , rotational mechanism | -0.5085 | -1.4 | 0.0171782 |
| YHL040C | ARN1 | siderophore-iron transporter | -0.5083 | -1.4 | 0.0553052 |
| YJR047C | ANB1 | translation initiation factor | -0.5077 | -1.4 | 0.0265618 |
| YOR270C | VPH1 | hydrogen-transporting ATPase , rotational mechanism | -0.5073 | -1.4 | 0.328298 |
| YIL018W | RPL2B | structural constituent of ribosome | -0.5068 | -1.4 | 0.045769 |
| YOL038W | PRE6 | endopeptidase | -0.5058 | -1.4 | 0.0178029 |
| YGR037C | ACB1 | long-chain fatty acid transporter | -0.5038 | -1.4 | 0.121755 |
| YBL113C | | helicase | -0.5020 | -1.4 | 0.0549479 |
| YDR134C | | unknown | -0.5003 | -1.4 | 0.0989924 |
| YHR027C | RPN1 | endopeptidase | -0.4963 | -1.4 | 0.0531673 |
| YDR300C | PRO1 | glutamate 5-kinase | -0.4957 | -1.4 | 0.0155524 |
| YER072W | VTC1 | unknown | -0.4933 | -1.4 | 0.100826 |
| YKR074W | | unknown | -0.4923 | -1.4 | 0.169534 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YKL107W | | unknown | -0.4913 | -1.4 | 0.471983 |
| YBR095C | RXT2 | unknown | -0.4863 | -1.4 | 0.0195059 |
| YNL209W | SSB2 | ATPase * | -0.4847 | -1.4 | 0.042555 |
| YNL151C | RPC31 | DNA-directed RNA polymerase | -0.4830 | -1.4 | 0.0393664 |
| YGR235C | | unknown | -0.4823 | -1.4 | 0.0581392 |
| YDR542W | PAU10 | unknown | -0.4807 | -1.4 | 0.0064604 |
| YLR314C | CDC3 | structural constituent of cytoskeleton* | -0.4807 | -1.4 | 0.27661 |
| YBR080C | SEC18 | ATPase | -0.4805 | -1.4 | 0.0162414 |
| YOR027W | STI1 | unfolded protein binding | -0.4797 | -1.4 | 0.0921955 |
| YJL161W | FMP33 | unknown | -0.4790 | -1.4 | 0.1197 |
| YLL050C | COF1 | protein binding | -0.4773 | -1.4 | 0.0410656 |
| YIL047C | SYG1 | unknown | -0.4772 | -1.4 | 0.219427 |
| YJL159W | HSP150 | structural constituent of cell wall | -0.4762 | -1.4 | 0.0321015 |
| YNL015W | PBI2 | endopeptidase inhibitor | -0.4752 | -1.4 | 0.648855 |
| YJL121C | RPE1 | ribulose-phosphate 3-epimerase | -0.4735 | -1.4 | 0.0205524 |
| YOR198C | BFR1 | RNA binding | -0.4702 | -1.4 | 0.0099157 |
| YMR271C | URA10 | orotate phosphoribosyltransferase | -0.4698 | -1.4 | 0.052667 |
| YJL210W | PEX2 | protein binding | -0.4695 | -1.4 | 0.0126098 |
| YDR063W | | unknown | -0.4670 | -1.4 | 0.13161 |
| YIL147C | SLN1 | osmosensor * | -0.4657 | -1.4 | 0.0022519 |
| YPL091W | GLR1 | glutathione-disulfide reductase | -0.4633 | -1.4 | 0.0355704 |
| YDR028C | REG1 | protein phosphatase type 1 regulator | -0.4623 | -1.4 | 0.025451 |
| YHR001W | OSH7 | oxysterol binding | -0.4612 | -1.4 | 0.0819638 |
| YDR512C | EMI1 | unknown | -0.4607 | -1.4 | 0.106574 |
| YIR032C | DAL3 | ureidoglycolate hydrolase | -0.4607 | -1.4 | 0.0464308 |
| YKR096W | | unknown | -0.4577 | -1.4 | 0.0077315 |
| YFR039C | | unknown | -0.4562 | -1.4 | 0.180178 |
| YER177W | BMH1 | protein binding | -0.4548 | -1.4 | 0.0616217 |
| YLR258W | GSY2 | glycogen (starch) synthase | -0.4548 | -1.4 | 0.253974 |
| YAL067C | SEO1 | transporter | -0.4523 | -1.4 | 0.0732858 |
| YBR180W | DTR1 | multidrug transporter | -0.4467 | -1.4 | 0.0440688 |
| YDR321W | ASP1 | asparaginase | -0.4465 | -1.4 | 0.0151755 |
| YFR004W | RPN11 | endopeptidase | -0.4443 | -1.4 | 0.0214781 |
| YIL176C | PAU14 | unknown | -0.4425 | -1.4 | 0.0757757 |
| YEL047C | | fumarate reductase (NADH) | -0.4418 | -1.4 | 0.0369753 |
| YBR224W | | | -0.4417 | -1.4 | 0.122312 |
| YOR196C | LIP5 | lipoic acid synthase | -0.4412 | -1.4 | 0.132005 |
| YJL216C | | alpha-galactosidase * | -0.4408 | -1.4 | 0.0190883 |
| YFR033C | QCR6 | ubiquinol-cytochrome-c reductase | -0.4407 | -1.4 | 0.0155085 |
| YPL015C | HST2 | NAD-dependent histone deacetylase | -0.4407 | -1.4 | 0.0033929 |
| YPR100W | MRPL51 | structural constituent of ribosome | -0.4397 | -1.4 | 0.0499669 |
| YAL063C | FLO9 | mannose binding | -0.4392 | -1.4 | 0.0452028 |
| YDR454C | GUK1 | guanylate kinase | -0.4388 | -1.4 | 0.0526079 |
| YNL131W | TOM22 | protein transporter | -0.4382 | -1.4 | 0.0659353 |
| YLR193C | | unknown | -0.4380 | -1.4 | 0.0005422 |
| YBR234C | ARC40 | structural constituent of cytoskeleton | -0.4370 | -1.4 | 0.0397934 |
| YOR141C | ARP8 | unknown | -0.4368 | -1.4 | 0.025171 |
| YPL234C | TFP3 | hydrogen-transporting ATPase , rotational mechanism | -0.4347 | -1.4 | 0.182705 |
| YCRX17W | | | -0.4318 | -1.3 | 0.146365 |
| YBR036C | CSG2 | enzyme regulator | -0.4315 | -1.3 | 0.108648 |
| YNL121C | TOM70 | protein transporter | -0.4307 | -1.3 | 0.0039095 |
| YEL044W | IES6 | unknown | -0.4287 | -1.3 | 0.0572575 |
| YPL037C | EGD1 | unfolded protein binding | -0.4283 | -1.3 | 0.0148963 |
| YGL210W | YPT32 | GTPase | -0.4253 | -1.3 | 0.112283 |
| YLR429W | CRN1 | protein binding | -0.4247 | -1.3 | 0.0868638 |
| YJL223C | PAU1 | unknown | -0.4245 | -1.3 | 0.0038577 |
| YJR017C | ESS1 | peptidyl-prolyl cis-trans isomerase | -0.4233 | -1.3 | 0.0522744 |
| YJL158C | CIS3 | structural constituent of cell wall | -0.4225 | -1.3 | 0.32418 |
| YBR218C | PYC2 | pyruvate carboxylase | -0.4223 | -1.3 | 0.155177 |
| YML008C | ERG6 | sterol 24-C-methyltransferase | -0.4180 | -1.3 | 0.0140024 |
| YDR272W | GLO2 | hydroxyacylglutathione hydrolase | -0.4172 | -1.3 | 0.0705474 |
| YOL030W | GAS5 | 1,3-beta-glucanosyltransferase | -0.4165 | -1.3 | 0.0530601 |
| YML055W | SPC2 | protein binding | -0.4152 | -1.3 | 0.0367222 |
| YJL112W | MDV1 | unknown | -0.4145 | -1.3 | 0.0023037 |
| YGR255C | COQ6 | ubiquinone biosynthesis monooxygenase | -0.4122 | -1.3 | 0.0146451 |
| YLR149C | | unknown | -0.4117 | -1.3 | 0.0187874 |
| YDL220C | CDC13 | single-stranded DNA binding | -0.4092 | -1.3 | 0.204353 |
| YOR332W | VMA4 | hydrogen-transporting ATPase , rotational mechanism | -0.4080 | -1.3 | 0.0298209 |
| YDR494W | RSM28 | structural constituent of ribosome | -0.4078 | -1.3 | 0.231802 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YJL073W | JEM1 | unfolded protein binding | -0.4065 | -1.3 | 0.0172061 |
| YMR008C | PLB1 | lysophospholipase | -0.4058 | -1.3 | 0.0436741 |
| YPR188C | MLC2 | myosin II binding | -0.4055 | -1.3 | 0.15973 |
| YAL049C | | unknown | -0.4048 | -1.3 | 0.0154981 |
| YCL043C | PDII | protein disulfide isomerase | -0.4033 | -1.3 | 0.0204957 |
| YCL017C | NFS1 | cystathionine gamma-lyase * | -0.4025 | -1.3 | 0.0820098 |
| YNL079C | TPM1 | actin lateral binding | -0.4023 | -1.3 | 0.0002213 |
| YER174C | GRX4 | thiol-disulfide exchange intermediate | -0.4018 | -1.3 | 0.0834665 |
| YMR027W | | unknown | -0.4007 | -1.3 | 0.0303282 |
| YOR008C | SLG1 | transmembrane receptor | -0.4007 | -1.3 | 0.12183 |
| YBR047W | FMP23 | unknown | -0.4002 | -1.3 | 0.0768826 |
| YCR077C | PAT1 | unknown | -0.3997 | -1.3 | 0.152069 |
| YJL206C-A | | | -0.3988 | -1.3 | 0.27333 |
| YDR161W | | unknown | -0.3982 | -1.3 | 0.0124273 |
| YDR027C | VPS54 | unknown | -0.3968 | -1.3 | 0.413022 |
| YFR015C | GSY1 | glycogen (starch) synthase | -0.3958 | -1.3 | 0.295452 |
| YPL087W | YDC1 | ceramidase | -0.3950 | -1.3 | 0.0612491 |
| YAL068C | PAU8 | unknown | -0.3932 | -1.3 | 0.0152362 |
| YGL003C | CDH1 | enzyme activator | -0.3932 | -1.3 | 0.0087865 |
| YBR197C | | unknown | -0.3928 | -1.3 | 0.0273176 |
| YIL150C | MCM10 | chromatin binding | -0.3928 | -1.3 | 0.193376 |
| YNL022C | | unknown | -0.3922 | -1.3 | 0.0431623 |
| YCL028W | RNQ1 | unknown | -0.3903 | -1.3 | 0.0402925 |
| YGR044C | RME1 | specific transcriptional repressor | -0.3897 | -1.3 | 0.0137378 |
| YJR133W | XPT1 | xanthine phosphoribosyltransferase | -0.3897 | -1.3 | 0.0138571 |
| YOL120C | RPL18A | structural constituent of ribosome | -0.3892 | -1.3 | 0.101527 |
| YJL105W | SET4 | unknown | -0.3853 | -1.3 | 0.0464239 |
| YIL094C | LYS12 | homoisocitrate dehydrogenase | -0.3838 | -1.3 | 0.068896 |
| YBR158W | AMN1 | protein binding | -0.3830 | -1.3 | 0.136506 |
| YHR121W | LSM12 | RNA binding | -0.3812 | -1.3 | 0.0055828 |
| YCR102C | | unknown | -0.3795 | -1.3 | 0.0525116 |
| YDR201W | SPC19 | structural constituent of cytoskeleton | -0.3787 | -1.3 | 0.0574842 |
| YPR023C | EAF3 | histone acetyltransferase | -0.3763 | -1.3 | 0.0535381 |
| YMR260C | TIF11 | translation initiation factor | -0.3757 | -1.3 | 0.051849 |
| YBR082C | UBC4 | ubiquitin conjugating enzyme | -0.3750 | -1.3 | 0.101735 |
| YLR055C | SPT8 | transcription cofactor | -0.3750 | -1.3 | 0.0440691 |
| YIL156W | UBP7 | ubiquitin-specific protease | -0.3705 | -1.3 | 0.057431 |
| YIL087C | | unknown | -0.3703 | -1.3 | 0.073208 |
| YHR208W | BAT1 | branched-chain-amino-acid transaminase | -0.3693 | -1.3 | 0.0252525 |
| YBR006W | UGA2 | succinate-semialdehyde dehydrogenase [NAD(P)+] | -0.3687 | -1.3 | 0.102173 |
| YJR008W | | unknown | -0.3680 | -1.3 | 0.0023637 |
| YDR150W | NUM1 | tubulin binding | -0.3672 | -1.3 | 0.123314 |
| YBR269C | FMP21 | unknown | -0.3663 | -1.3 | 0.025059 |
| YHL028W | WSC4 | transmembrane receptor | -0.3620 | -1.3 | 0.373484 |
| YOR108W | LEU9 | 2-isopropylmalate synthase | -0.3613 | -1.3 | 0.278176 |
| YLR242C | ARV1 | unknown | -0.3607 | -1.3 | 0.0431188 |
| YIL109C | SEC24 | protein binding | -0.3532 | -1.3 | 0.0583009 |
| YCR049C | | | -0.3527 | -1.3 | 0.129834 |
| YMR325W | PAU19 | unknown | -0.3527 | -1.3 | 0.191408 |
| YER052C | HOM3 | aspartate kinase | -0.3517 | -1.3 | 0.0395801 |
| YDR109C | | kinase | -0.3502 | -1.3 | 0.0862926 |
| YER094C | PUP3 | endopeptidase | -0.3502 | -1.3 | 0.199199 |
| YJR064W | CCT5 | unfolded protein binding | -0.3500 | -1.3 | 0.189069 |
| YBL002W | HTB2 | DNA binding | -0.3490 | -1.3 | 0.0963136 |
| YGL112C | TAF6 | general RNA polymerase II transcription factor | -0.3475 | -1.3 | 0.0965273 |
| YGR181W | TIM13 | protein transporter | -0.3452 | -1.3 | 0.291063 |
| YEL002C | WBP1 | dolichyl-diphosphooligosaccharide-protein glycotransferase | -0.3450 | -1.3 | 0.020492 |
| YIL086C | | | -0.3447 | -1.3 | 0.0208902 |
| YJL025W | RRN7 | RNA polymerase I transcription factor | -0.3435 | -1.3 | 0.524392 |
| YLL013C | PUF3 | mRNA binding | -0.3425 | -1.3 | 0.0467786 |
| YOR362C | PRE10 | endopeptidase | -0.3385 | -1.3 | 0.183092 |
| YPL033C | | unknown | -0.3342 | -1.3 | 0.127051 |
| YJL055W | | unknown | -0.3335 | -1.3 | 0.116933 |
| YJL202C | | | -0.3333 | -1.3 | 0.275421 |
| YER022W | SRB4 | RNA polymerase II transcription mediator | -0.3328 | -1.3 | 0.296652 |
| YHR009C | | unknown | -0.3313 | -1.3 | 0.013602 |
| YER066C-A | | | -0.3308 | -1.3 | 0.280224 |
| YDL135C | RDI1 | signal transducer | -0.3278 | -1.3 | 0.0184586 |
| YJL153C | INO1 | inositol-3-phosphate synthase | -0.3277 | -1.3 | 0.0728575 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YHR026W | PPA1 | hydrogen-transporting ATPase , rotational mechanism | -0.3270 | -1.3 | 0.01885 |
| YDR079W | PET100 | unfolded protein binding | -0.3248 | -1.3 | 0.12494 |
| YPR004C | | electron carrier | -0.3232 | -1.3 | 0.0088409 |
| YDR033W | MRH1 | unknown | -0.3228 | -1.3 | 0.129756 |
| YJL070C | | unknown | -0.3210 | -1.2 | 0.0965442 |
| YPR019W | CDC54 | chromatin binding | -0.3208 | -1.2 | 0.0691866 |
| YDR394W | RPT3 | ATPase * | -0.3200 | -1.2 | 0.0142985 |
| YIL005W | EPS1 | protein disulfide isomerase | -0.3187 | -1.2 | 0.0147456 |
| YBR132C | AGP2 | amino acid transporter | -0.3160 | -1.2 | 0.0530281 |
| YEL046C | GLY1 | threonine aldolase | -0.3110 | -1.2 | 0.0473239 |
| YDR133C | | | -0.3107 | -1.2 | 0.245479 |
| YCR087W | | | -0.3103 | -1.2 | 0.328557 |
| YCR104W | PAU3 | unknown | -0.3093 | -1.2 | 0.463135 |
| YDL086W | | carboxymethylenebutenolidase | -0.3058 | -1.2 | 0.108362 |
| YDR320C | SWA2 | protein binding | -0.3053 | -1.2 | 0.129512 |
| YBR090C | | unknown | -0.3045 | -1.2 | 0.0116947 |
| YHR217C | | | -0.3043 | -1.2 | 0.0116195 |
| YGL073W | HSF1 | transcription factor | -0.3038 | -1.2 | 0.14289 |
| YMR311C | GLC8 | enzyme activator | -0.3027 | -1.2 | 0.0116322 |
| YGL147C | RPL9A | structural constituent of ribosome | -0.3020 | -1.2 | 0.121024 |
| YDR059C | UBC5 | ubiquitin conjugating enzyme | -0.3007 | -1.2 | 0.128747 |
| YLR048W | RPS0B | structural constituent of ribosome | -0.2995 | -1.2 | 0.018032 |
| YIL098C | FMC1 | unknown | -0.2973 | -1.2 | 0.236249 |
| YLR102C | APC9 | protein binding | -0.2965 | -1.2 | 0.213915 |
| YBR227C | MCX1 | unfolded protein binding | -0.2962 | -1.2 | 0.204366 |
| YDR255C | RMD5 | unknown | -0.2952 | -1.2 | 0.0875604 |
| YNL138W | SRV2 | cytoskeletal protein binding | -0.2920 | -1.2 | 0.100936 |
| YOR383C | FIT3 | unknown | -0.2915 | -1.2 | 0.411951 |
| YDL178W | DLD2 | actin binding | -0.2910 | -1.2 | 0.454079 |
| YPR090W | | | -0.2907 | -1.2 | 0.0998667 |
| YAL005C | SSA1 | ATPase * | -0.2905 | -1.2 | 0.286351 |
| YBR222C | PCS60 | AMP binding | -0.2898 | -1.2 | 0.0200898 |
| YGR008C | STF2 | unknown | -0.2895 | -1.2 | 0.0064264 |
| YPL094C | SEC62 | protein binding | -0.2893 | -1.2 | 0.095558 |
| YIL162W | SUC2 | beta-fructofuranosidase | -0.2890 | -1.2 | 0.113941 |
| YFR017C | | unknown | -0.2887 | -1.2 | 0.189414 |
| YGR135W | PRE9 | endopeptidase | -0.2868 | -1.2 | 0.201021 |
| YOR374W | ALD4 | aldehyde dehydrogenase (NAD) | -0.2868 | -1.2 | 0.0791755 |
| YGR232W | NAS6 | unknown | -0.2867 | -1.2 | 0.0187189 |
| YLR299W | ECM38 | protein-glutamine gamma-glutamyltransferase | -0.2863 | -1.2 | 0.187261 |
| YMR100W | MUB1 | unknown | -0.2858 | -1.2 | 0.0711819 |
| YMR196W | | unknown | -0.2837 | -1.2 | 0.277547 |
| YNL268W | LYP1 | basic amino acid transporter | -0.2835 | -1.2 | 0.426689 |
| YGR209C | TRX2 | thiol-disulfide exchange intermediate | -0.2827 | -1.2 | 0.189644 |
| YNL104C | LEU4 | 2-isopropylmalate synthase | -0.2812 | -1.2 | 0.0032684 |
| YER057C | HMF1 | unknown | -0.2803 | -1.2 | 0.281469 |
| YOL078W | AVO1 | unknown | -0.2793 | -1.2 | 0.0549599 |
| YKR051W | | unknown | -0.2775 | -1.2 | 0.0474545 |
| YPL106C | SSE1 | ATP binding | -0.2773 | -1.2 | 0.126718 |
| YFL004W | VTC2 | unknown | -0.2747 | -1.2 | 0.195109 |
| YBR245C | ISW1 | ATPase | -0.2737 | -1.2 | 0.05567 |
| YDL130W | RPP1B | structural constituent of ribosome | -0.2732 | -1.2 | 0.106492 |
| YGL056C | SDS23 | unknown | -0.2723 | -1.2 | 0.0126981 |
| YBR262C | FMP51 | unknown | -0.2708 | -1.2 | 0.0120791 |
| YJL166W | QCR8 | ubiquinol-cytochrome-c reductase | -0.2697 | -1.2 | 0.0387817 |
| YLR204W | QRI5 | unknown | -0.2695 | -1.2 | 0.199011 |
| YPL190C | NAB3 | poly(A) binding | -0.2685 | -1.2 | 0.131304 |
| YDL184C | RPL41A | structural constituent of ribosome | -0.2672 | -1.2 | 0.191252 |
| YBR164C | ARL1 | GTPase | -0.2667 | -1.2 | 0.160548 |
| YJL167W | ERG20 | dimethylallyltranstransferase * | -0.2667 | -1.2 | 0.0354019 |
| YBR102C | EXO84 | protein binding | -0.2665 | -1.2 | 0.139396 |
| YCL051W | LRE1 | transcription regulator * | -0.2625 | -1.2 | 0.54067 |
| YJL115W | ASF1 | transcription regulator * | -0.2602 | -1.2 | 0.370579 |
| YPL177C | CUP9 | specific RNA polymerase II transcription factor | -0.2598 | -1.2 | 0.0465961 |
| YGR193C | PDX1 | protein binding | -0.2577 | -1.2 | 0.12853 |
| YML100W | TSL1 | enzyme regulator * | -0.2568 | -1.2 | 0.370253 |
| YGR189C | CRH1 | unknown | -0.2565 | -1.2 | 0.137615 |
| YGL108C | | unknown | -0.2552 | -1.2 | 0.132556 |
| YDR329C | PEX3 | protein binding | -0.2523 | -1.2 | 0.125236 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YMR043W | MCM1 | DNA binding | -0.2513 | -1.2 | 0.0104057 |
| YPR002W | PDH1 | unknown | -0.2512 | -1.2 | 0.0285461 |
| YLR340W | RPP0 | structural constituent of ribosome | -0.2507 | -1.2 | 0.12604 |
| YIL138C | TPM2 | actin lateral binding | -0.2503 | -1.2 | 0.136127 |
| YML103C | NUP188 | structural molecule | -0.2503 | -1.2 | 0.47276 |
| YCL037C | SRO9 | RNA binding | -0.2495 | -1.2 | 0.670089 |
| YDR091C | RLI1 | ATPase * | -0.2487 | -1.2 | 0.0577111 |
| YIL160C | POT1 | acetyl-CoA C-acyltransferase | -0.2480 | -1.2 | 0.290992 |
| YBL027W | RPL19B | structural constituent of ribosome | -0.2478 | -1.2 | 0.288063 |
| YPL040C | ISM1 | isoleucine-tRNA ligase | -0.2468 | -1.2 | 0.404087 |
| YBR202W | CDC47 | chromatin binding | -0.2465 | -1.2 | 0.0574305 |
| YIL084C | SDS3 | protein binding | -0.2452 | -1.2 | 0.196977 |
| YGL019W | CKB1 | protein kinase | -0.2447 | -1.2 | 0.184944 |
| YNL160W | YGP1 | unknown | -0.2443 | -1.2 | 0.296927 |
| YEL027W | CUP5 | hydrogen ion transporter | -0.2412 | -1.2 | 0.0395769 |
| YLR043C | TRX1 | thiol-disulfide exchange intermediate | -0.2412 | -1.2 | 0.368864 |
| YIL088C | AVT7 | transporter | -0.2407 | -1.2 | 0.151876 |
| YKL058W | TOA2 | general RNA polymerase II transcription factor | -0.2397 | -1.2 | 0.181891 |
| YPL085W | SEC16 | structural molecule | -0.2387 | -1.2 | 0.711948 |
| YBR184W | | unknown | -0.2375 | -1.2 | 0.540746 |
| YDR080W | VPS41 | Rab guanyl-nucleotide exchange factor | -0.2368 | -1.2 | 0.0940768 |
| YBR147W | | unknown | -0.2363 | -1.2 | 0.313067 |
| YCL027W | FUS1 | unknown | -0.2353 | -1.2 | 0.0037438 |
| YCL056C | | unknown | -0.2347 | -1.2 | 0.204913 |
| YBR216C | YBP1 | unknown | -0.2323 | -1.2 | 0.185157 |
| YGL219C | MDM34 | unknown | -0.2323 | -1.2 | 0.0003069 |
| YHR129C | ARP1 | structural constituent of cytoskeleton | -0.2320 | -1.2 | 0.687546 |
| YPR108W | RPN7 | structural molecule | -0.2313 | -1.2 | 0.0626768 |
| YER002W | NOP16 | unknown | -0.2297 | -1.2 | 0.0468075 |
| YDR115W | | structural constituent of ribosome | -0.2267 | -1.2 | 0.118588 |
| YLR247C | | helicase | -0.2267 | -1.2 | 0.120682 |
| YER035W | EDC2 | RNA binding | -0.2262 | -1.2 | 0.0925406 |
| YDR427W | RPN9 | structural molecule | -0.2262 | -1.2 | 0.0070353 |
| YJL123C | | unknown | -0.2245 | -1.2 | 0.179128 |
| YAL013W | DEP1 | transcription regulator | -0.2237 | -1.2 | 0.265207 |
| YHR131C | | unknown | -0.2233 | -1.2 | 0.203873 |
| YER119C | AVT6 | amino acid transporter | -0.2232 | -1.2 | 0.0782718 |
| YOR239W | ABP140 | S-adenosylmethionine-dependent methyltransferase * | -0.2205 | -1.2 | 0.0022359 |
| YGL104C | VPS73 | unknown | -0.2202 | -1.2 | 0.480298 |
| YBR237W | PRP5 | RNA splicing factor , transesterification mechanism* | -0.2197 | -1.2 | 0.0681174 |
| YNL125C | ESBP6 | transporter | -0.2195 | -1.2 | 0.324223 |
| YBR017C | KAP104 | nuclear localization sequence binding | -0.2192 | -1.2 | 0.23811 |
| YHR215W | PHO12 | acid phosphatase | -0.2125 | -1.2 | 0.02025 |
| YJL209W | CBP1 | mRNA binding | -0.2113 | -1.2 | 0.451795 |
| YCL033C | | protein-methionine-R-oxide reductase | -0.2110 | -1.2 | 0.0413544 |
| YLR355C | ILV5 | ketol-acid reductoisomerase | -0.2095 | -1.2 | 0.176056 |
| YGR284C | ERV29 | unknown | -0.2050 | -1.2 | 0.0496765 |
| YMR295C | | unknown | -0.2050 | -1.2 | 0.182676 |
| YER158C | | unknown | -0.2048 | -1.2 | 0.256225 |
| YEL051W | VMA8 | hydrogen-transporting ATPase , rotational mechanism | -0.2042 | -1.2 | 0.0897327 |
| YMR135C | GID8 | unknown | -0.2032 | -1.2 | 0.419044 |
| YDR169C | STB3 | unknown | -0.2025 | -1.2 | 0.389068 |
| YMR233W | | unknown | -0.2003 | -1.1 | 0.123669 |
| YBR261C | | S-adenosylmethionine-dependent methyltransferase | -0.1990 | -1.1 | 0.323767 |
| YLR216C | CPR6 | unfolded protein binding | -0.1990 | -1.1 | 0.121774 |
| YER063W | THO1 | unknown | -0.1972 | -1.1 | 0.115254 |
| YDR116C | MRPL1 | structural constituent of ribosome | -0.1962 | -1.1 | 0.217686 |
| YAL003W | EFB1 | translation elongation factor | -0.1957 | -1.1 | 0.18672 |
| YDR481C | PHO8 | alkaline phosphatase | -0.1948 | -1.1 | 0.17067 |
| YGL031C | RPL24A | structural constituent of ribosome* | -0.1920 | -1.1 | 0.150615 |
| YHR025W | THR1 | homoserine kinase | -0.1917 | -1.1 | 0.112021 |
| YMR264W | CUE1 | protein binding | -0.1915 | -1.1 | 0.357225 |
| YDR232W | HEM1 | 5-aminolevulinate synthase | -0.1912 | -1.1 | 0.0846545 |
| YKL204W | EAP1 | eukaryotic initiation factor 4E binding | -0.1885 | -1.1 | 0.372457 |
| YOR197W | MCA1 | caspase | -0.1880 | -1.1 | 0.0302436 |
| YEL053C | MAK10 | peptide alpha-N-acetyltransferase | -0.1852 | -1.1 | 0.297628 |
| YPL093W | NOG1 | GTP binding | -0.1830 | -1.1 | 0.494304 |
| YCR026C | NPP1 | nucleoside-triphosphatase * | -0.1822 | -1.1 | 0.329519 |
| YDR517W | GRH1 | unknown | -0.1822 | -1.1 | 0.240009 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YGL059W | | protein kinase | -0.1818 | -1.1 | 0.0240171 |
| YHR200W | RPN10 | endopeptidase | -0.1792 | -1.1 | 0.328357 |
| YPL105C | | unknown | -0.1782 | -1.1 | 0.13231 |
| YGR180C | RNR4 | ribonucleoside-diphosphate reductase | -0.1768 | -1.1 | 0.23795 |
| YBR137W | | unknown | -0.1752 | -1.1 | 0.0905536 |
| YCL001W | RER1 | unknown | -0.1748 | -1.1 | 0.396328 |
| YJL124C | LSM1 | RNA binding | -0.1743 | -1.1 | 0.492226 |
| YPL271W | ATP15 | hydrogen-transporting ATP synthase , rotational mechanism | -0.1737 | -1.1 | 0.666933 |
| YLR436C | ECM30 | unknown | -0.1733 | -1.1 | 0.663404 |
| YJR005W | APL1 | unknown | -0.1725 | -1.1 | 0.617463 |
| YNL067W | RPL9B | structural constituent of ribosome | -0.1718 | -1.1 | 0.259672 |
| YPL259C | APM1 | clathrin binding | -0.1708 | -1.1 | 0.0661125 |
| YJL095W | BCK1 | MAP kinase | -0.1700 | -1.1 | 0.0708709 |
| YBR151W | APD1 | unknown | -0.1675 | -1.1 | 0.300384 |
| YJL026W | RNR2 | ribonucleoside-diphosphate reductase | -0.1673 | -1.1 | 0.210331 |
| YNL102W | POL1 | alpha DNA polymerase | -0.1663 | -1.1 | 0.285585 |
| YBR250W | | unknown | -0.1653 | -1.1 | 0.286503 |
| YFL010C | WWM1 | unknown | -0.1650 | -1.1 | 0.0138209 |
| YMR241W | YHM2 | DNA binding | -0.1632 | -1.1 | 0.019435 |
| YIR014W | | unknown | -0.1625 | -1.1 | 0.308478 |
| YJL129C | TRK1 | potassium ion transporter | -0.1607 | -1.1 | 0.44731 |
| YLR350W | ORM2 | unknown | -0.1598 | -1.1 | 0.301149 |
| YNL241C | ZWF1 | glucose-6-phosphate 1-dehydrogenase | -0.1588 | -1.1 | 0.0377215 |
| YLR064W | | unknown | -0.1582 | -1.1 | 0.482771 |
| YER060W-A | FCY22 | cytosine-purine permease | -0.1567 | -1.1 | 0.43332 |
| YKR049C | FMP46 | oxidoreductase | -0.1552 | -1.1 | 0.493468 |
| YIR007W | | unknown | -0.1530 | -1.1 | 0.235529 |
| YNL245C | CWC25 | unknown | -0.1515 | -1.1 | 0.111736 |
| YBR118W | TEF2 | translation elongation factor | -0.1512 | -1.1 | 0.512129 |
| YBR149W | ARA1 | aldo-keto reductase * | -0.1512 | -1.1 | 0.693718 |
| YDL173W | | unknown | -0.1507 | -1.1 | 0.169142 |
| YPL203W | TPK2 | protein serine/threonine kinase | -0.1505 | -1.1 | 0.136663 |
| YBR101C | FES1 | adenyl-nucleotide exchange factor | -0.1502 | -1.1 | 0.286266 |
| YDL014W | NOP1 | methyltransferase | -0.1498 | -1.1 | 0.165326 |
| YHR089C | GAR1 | RNA binding | -0.1497 | -1.1 | 0.570321 |
| YGL150C | INO80 | ATPase * | -0.1493 | -1.1 | 0.215053 |
| YCL052C | PBN1 | mannosyltransferase | -0.1482 | -1.1 | 0.402806 |
| YER142C | MAG1 | alkylbase DNA N-glycosylase | -0.1482 | -1.1 | 0.523804 |
| YOR261C | RPN8 | unknown | -0.1478 | -1.1 | 0.413071 |
| YPL240C | HSP82 | ATPase , coupled | -0.1437 | -1.1 | 0.140038 |
| YHR194W | MDM31 | unknown | -0.1422 | -1.1 | 0.26376 |
| YIL154C | IMP2' | transcription coactivator | -0.1415 | -1.1 | 0.0942467 |
| YIL062C | ARC15 | structural molecule * | -0.1410 | -1.1 | 0.200793 |
| YJL156C | SSY5 | amino acid binding | -0.1393 | -1.1 | 0.484104 |
| YJR127C | RSF2 | transcription factor | -0.1392 | -1.1 | 0.130358 |
| YEL012W | UBC8 | ubiquitin conjugating enzyme | -0.1370 | -1.1 | 0.192647 |
| YLR190W | MMR1 | unknown | -0.1348 | -1.1 | 0.107811 |
| YLR437C | | unknown | -0.1348 | -1.1 | 0.444345 |
| YGR155W | CYS4 | cystathionine beta-synthase | -0.1345 | -1.1 | 0.184863 |
| YJL137C | GLG2 | glycogenin glucosyltransferase | -0.1342 | -1.1 | 0.292176 |
| YKR062W | TFA2 | general RNA polymerase II transcription factor | -0.1337 | -1.1 | 0.121212 |
| YJR104C | SOD1 | copper, zinc superoxide dismutase | -0.1328 | -1.1 | 0.229208 |
| YOR327C | SNC2 | v-SNARE | -0.1327 | -1.1 | 0.240595 |
| YDR156W | RPA14 | DNA-directed RNA polymerase | -0.1325 | -1.1 | 0.283442 |
| YKL064W | MNR2 | magnesium ion transporter | -0.1312 | -1.1 | 0.334449 |
| YKR059W | TIF1 | translation initiation factor | -0.1308 | -1.1 | 0.399465 |
| YJL151C | SNA3 | unknown | -0.1260 | -1.1 | 0.349272 |
| YAL039C | CYC3 | holocytochrome-c synthase | -0.1255 | -1.1 | 0.226222 |
| YBR240C | THI2 | transcriptional activator | -0.1247 | -1.1 | 0.184372 |
| YJR105W | ADO1 | adenosine kinase | -0.1243 | -1.1 | 0.258849 |
| YAL061W | | oxidoreductase | -0.1230 | -1.1 | 0.396 |
| YLR005W | SSL1 | general RNA polymerase II transcription factor | -0.1218 | -1.1 | 0.716853 |
| YGL084C | GUP1 | O-acyltransferase | -0.1203 | -1.1 | 0.117724 |
| YGR236C | SPG1 | unknown | -0.1203 | -1.1 | 0.0253991 |
| YBR053C | | unknown | -0.1200 | -1.1 | 0.0275542 |
| YFL018W-A | | | -0.1197 | -1.1 | 0.295279 |
| YAL011W | SWC3 | unknown | -0.1187 | -1.1 | 0.589352 |
| YBR156C | SLI15 | protein kinase | -0.1182 | -1.1 | 0.238679 |
| YML048W | GSF2 | unknown | -0.1178 | -1.1 | 0.41546 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YDR128W | | unknown | -0.1167 | -1.1 | 0.19689 |
| YBL050W | SEC17 | soluble NSF attachment protein | -0.1165 | -1.1 | 0.572066 |
| YDR378C | LSM6 | RNA binding | -0.1165 | -1.1 | 0.192762 |
| YLR192C | HCR1 | translation initiation factor * | -0.1165 | -1.1 | 0.713732 |
| YHR096C | HXT5 | glucose transporter | -0.1163 | -1.1 | 0.674733 |
| YIL046W | MET30 | protein binding | -0.1155 | -1.1 | 0.652273 |
| YBR204C | | serine hydrolase | -0.1143 | -1.1 | 0.615932 |
| YGR285C | ZUO1 | unfolded protein binding | -0.1133 | -1.1 | 0.136678 |
| YBR059C | AKL1 | protein kinase | -0.1132 | -1.1 | 0.465755 |
| YHR158C | KEL1 | unknown | -0.1123 | -1.1 | 0.896318 |
| YDR382W | RPP2B | structural constituent of ribosome | -0.1120 | -1.1 | 0.190768 |
| YGL011C | SCL1 | endopeptidase | -0.1110 | -1.1 | 0.507073 |
| YHR179W | OYE2 | NADPH dehydrogenase | -0.1105 | -1.1 | 0.317048 |
| YIL050W | PCL7 | cyclin-dependent protein kinase | -0.1103 | -1.1 | 0.363737 |
| YPR062W | FCY1 | cytosine deaminase | -0.1088 | -1.1 | 0.0745763 |
| YAL035W | FUN12 | GTPase * | -0.1080 | -1.1 | 0.326953 |
| YDL156W | | unknown | -0.1058 | -1.1 | 0.216533 |
| YIL142W | CCT2 | unfolded protein binding | -0.1055 | -1.1 | 0.202652 |
| YDR179W-A | | unknown | -0.1052 | -1.1 | 0.131808 |
| YJL138C | TIF2 | translation initiation factor * | -0.1033 | -1.1 | 0.319223 |
| YNL208W | | unknown | -0.1025 | -1.1 | 0.523182 |
| YOR185C | GSP2 | GTPase | -0.1018 | -1.1 | 0.0915285 |
| YKL013C | ARC19 | structural molecule | -0.1008 | -1.1 | 0.271515 |
| YBR238C | | unknown | -0.0993 | -1.1 | 0.267861 |
| YGL197W | MDS3 | unknown | -0.0983 | -1.1 | 0.849497 |
| YBR159W | IFA38 | ketoreductase | -0.0977 | -1.1 | 0.844389 |
| YOR286W | FMP31 | unknown | -0.0977 | -1.1 | 0.476641 |
| YPL090C | RPS6A | structural constituent of ribosome | -0.0935 | -1.1 | 0.235743 |
| YOR015W | | | -0.0920 | -1.1 | 0.656495 |
| YJL101C | GSH1 | glutamate-cysteine ligase | -0.0905 | -1.1 | 0.164251 |
| YBR255W | | unknown | -0.0897 | -1.1 | 0.576046 |
| YJR085C | | unknown | -0.0890 | -1.1 | 0.556205 |
| YHR192W | | unknown | -0.0882 | -1.1 | 0.570074 |
| YBR247C | ENP1 | snoRNA binding | -0.0872 | -1.1 | 0.773009 |
| YNL056W | OCA2 | unknown | -0.0872 | -1.1 | 0.136971 |
| YBR200W | BEM1 | protein binding | -0.0868 | -1.1 | 0.371361 |
| YMR302C | YME2 | exonuclease | -0.0868 | -1.1 | 0.684504 |
| YCL008C | STP22 | protein binding | -0.0863 | -1.1 | 0.32417 |
| YOL016C | CMK2 | calcium- and calmodulin-dependent protein kinase | -0.0858 | -1.1 | 0.692862 |
| YPL111W | CAR1 | zinc ion binding | -0.0852 | -1.1 | 0.406608 |
| YDR497C | ITR1 | myo-inositol transporter | -0.0847 | -1.1 | 0.415047 |
| YIL085C | KTR7 | mannosyltransferase | -0.0842 | -1.1 | 0.47342 |
| YJR024C | | unknown | -0.0820 | -1.1 | 0.195425 |
| YBR069C | TAT1 | amino acid transporter | -0.0765 | -1.1 | 0.686578 |
| YBR070C | ALG14 | N-acetylglucosaminyl diphosphodolichol N-acetylglucosaminyl | -0.0758 | -1.1 | 0.459492 |
| YOR123C | LEO1 | RNA polymerase II transcription elongation factor | -0.0752 | -1.1 | 0.13178 |
| YDL095W | PMT1 | dolichyl-phosphate-mannose-protein mannosyltransferase | -0.0738 | -1.1 | 0.483306 |
| YBR272C | HSM3 | unknown | -0.0737 | -1.1 | 0.69541 |
| YDR208W | MSS4 | 1-phosphatidylinositol 4-phosphate 5-kinase | -0.0698 | -1.0 | 0.123837 |
| YKR097W | PCK1 | phosphoenolpyruvate carboxykinase | -0.0695 | -1.0 | 0.394403 |
| YDR358W | GGA1 | ubiquitin binding | -0.0693 | -1.0 | 0.610278 |
| YCR018C | SRD1 | unknown | -0.0665 | -1.0 | 0.642644 |
| YJL113W | | | -0.0652 | -1.0 | 0.7457 |
| YJL221C | FSP2 | alpha-glucosidase | -0.0638 | -1.0 | 0.821865 |
| YCRX13W | | | -0.0627 | -1.0 | 0.153112 |
| YJL016W | | unknown | -0.0618 | -1.0 | 0.748095 |
| YBR223C | TDP1 | tyrosyl-DNA phosphodiesterase | -0.0590 | -1.0 | 0.829929 |
| YER067W | | unknown | -0.0588 | -1.0 | 0.656957 |
| YBR111C | YSA1 | phosphoribosyl-ATP diphosphatase | -0.0567 | -1.0 | 0.326224 |
| YHR193C | EGD2 | unfolded protein binding | -0.0562 | -1.0 | 0.813478 |
| YMR186W | HSC82 | unfolded protein binding | -0.0558 | -1.0 | 0.571664 |
| YKL100C | | unknown | -0.0547 | -1.0 | 0.83955 |
| YBR259W | | unknown | -0.0543 | -1.0 | 0.451349 |
| YDR056C | | unknown | -0.0533 | -1.0 | 0.367201 |
| YIL068C | SEC6 | protein binding | -0.0523 | -1.0 | 0.510106 |
| YBR252W | DUT1 | dUTP diphosphatase | -0.0513 | -1.0 | 0.757945 |
| YNL147W | LSM7 | RNA binding | -0.0510 | -1.0 | 0.402097 |
| YMR230W | RPS10B | structural constituent of ribosome | -0.0503 | -1.0 | 0.631628 |
| YOL032W | OPI10 | unknown | -0.0500 | -1.0 | 0.223904 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|----------|
| YBR057C | MUM2 | unknown | -0.0497 | -1.0 | 0.692495 |
| YNL126W | SPC98 | structural constituent of cytoskeleton | -0.0475 | -1.0 | 0.726938 |
| YAL029C | MYO4 | microfilament motor | -0.0472 | -1.0 | 0.916709 |
| YIL158W | | unknown | -0.0470 | -1.0 | 0.702289 |
| YGR118W | RPS23A | structural constituent of ribosome | -0.0463 | -1.0 | 0.400155 |
| YBR198C | TAF5 | general RNA polymerase II transcription factor | -0.0458 | -1.0 | 0.302126 |
| YIL130W | ASG1 | unknown | -0.0458 | -1.0 | 0.664978 |
| YKR075C | | unknown | -0.0440 | -1.0 | 0.7065 |
| YML063W | RPS1B | structural constituent of ribosome | -0.0437 | -1.0 | 0.814976 |
| YLR345W | | 6-phosphofructo-2-kinase | -0.0427 | -1.0 | 0.612659 |
| YMR020W | FMS1 | amine oxidase | -0.0423 | -1.0 | 0.61925 |
| YML073C | RPL6A | structural constituent of ribosome* | -0.0420 | -1.0 | 0.544518 |
| YBR170C | NPL4 | unknown | -0.0407 | -1.0 | 0.77877 |
| YNR021W | | unknown | -0.0372 | -1.0 | 0.601968 |
| YIL076W | SEC28 | unknown | -0.0362 | -1.0 | 0.851407 |
| YMR061W | RNA14 | RNA binding | -0.0357 | -1.0 | 0.747233 |
| YFL008W | SMC1 | ATPase * | -0.0355 | -1.0 | 0.807287 |
| YER054C | GIP2 | protein phosphatase regulator | -0.0343 | -1.0 | 0.738647 |
| YBR126C | TPS1 | alpha,alpha-trehalose-phosphate synthase (UDP-forming) | -0.0327 | -1.0 | 0.29786 |
| YBR302C | COS2 | unknown | -0.0327 | -1.0 | 0.870826 |
| YCRX02C | | | -0.0307 | -1.0 | 0.764771 |
| YAL002W | VPS8 | unknown | -0.0302 | -1.0 | 0.618338 |
| YIL041W | GVP36 | unknown | -0.0282 | -1.0 | 0.623516 |
| YBL095W | | unknown | -0.0275 | -1.0 | 0.823013 |
| YKL198C | PTK1 | protein kinase | -0.0273 | -1.0 | 0.8231 |
| YEL016C | NPP2 | nucleoside-triphosphatase * | -0.0267 | -1.0 | 0.727216 |
| YER117W | RPL23B | structural constituent of ribosome | -0.0263 | -1.0 | 0.730651 |
| YCL035C | GRX1 | thiol-disulfide exchange intermediate * | -0.0245 | -1.0 | 0.901073 |
| YGR138C | TPO2 | spermine transporter | -0.0225 | -1.0 | 0.888756 |
| YMR273C | ZDS1 | protein binding | -0.0192 | -1.0 | 0.886538 |
| YKL086W | SRX1 | oxidoreductase | -0.0190 | -1.0 | 0.971947 |
| YDL147W | RPN5 | unknown | -0.0173 | -1.0 | 0.846439 |
| YOR080W | DIA2 | DNA replication origin binding | -0.0165 | -1.0 | 0.702658 |
| YDL213C | NOP6 | unknown | -0.0145 | -1.0 | 0.881505 |
| YMR188C | MRPS17 | structural constituent of ribosome | -0.0140 | -1.0 | 0.930087 |
| YBR236C | ABD1 | mRNA (guanine-N7-)-methyltransferase | -0.0133 | -1.0 | 0.887663 |
| YER073W | ALD5 | 3-chloroallyl aldehyde dehydrogenase | -0.0123 | -1.0 | 0.899672 |
| YDR444W | | unknown | -0.0120 | -1.0 | 0.977128 |
| YBR207W | FTH1 | iron ion transporter | -0.0117 | -1.0 | 0.54532 |
| YKL116C | PRR1 | receptor signaling protein serine/threonine kinase | -0.0097 | -1.0 | 0.925469 |
| YKL211C | TRP3 | anthranilate synthase * | -0.0093 | -1.0 | 0.971409 |
| YLR381W | CTF3 | protein binding | -0.0093 | -1.0 | 0.948143 |
| YCR023C | | unknown | -0.0050 | -1.0 | 0.973556 |
| YER131W | RPS26B | structural constituent of ribosome | -0.0038 | -1.0 | 0.973754 |
| YLR160C | ASP3-4 | asparaginase | -0.0025 | -1.0 | 0.975087 |
| YBR210W | ERV15 | unknown | -0.0023 | -1.0 | 0.990589 |
| YHR120W | MSH1 | ATP binding | -0.0022 | -1.0 | 0.99433 |
| YGR196C | FYV8 | unknown | 0.0002 | 1.0 | 0.996927 |
| YLR447C | VMA6 | hydrogen-transporting ATPase , rotational mechanism | 0.0003 | 1.0 | 0.973757 |
| YBL066C | SEF1 | unknown | 0.0013 | 1.0 | 0.998359 |
| YOR375C | GDH1 | glutamate dehydrogenase (NADP+) | 0.0013 | 1.0 | 0.992311 |
| YBR188C | NTC20 | RNA splicing factor , transesterification mechanism | 0.0015 | 1.0 | 0.983857 |
| YDR514C | | unknown | 0.0022 | 1.0 | 0.992061 |
| YBR035C | PDX3 | pyridoxamine-phosphate oxidase | 0.0030 | 1.0 | 0.978258 |
| YAL030W | SNC1 | v-SNARE | 0.0035 | 1.0 | 0.817056 |
| YCL019W | | | 0.0037 | 1.0 | 0.983595 |
| YJL059W | YHC3 | basic amino acid transporter | 0.0038 | 1.0 | 0.94726 |
| YBR072W | HSP26 | unfolded protein binding | 0.0040 | 1.0 | 0.963297 |
| YOR063W | RPL3 | structural constituent of ribosome | 0.0040 | 1.0 | 0.915429 |
| YDR127W | ARO1 | 3-dehydroquinate dehydratase * | 0.0042 | 1.0 | 0.931692 |
| YJL111W | CCT7 | unfolded protein binding | 0.0045 | 1.0 | 0.941228 |
| YMR148W | | unknown | 0.0050 | 1.0 | 0.987813 |
| YOR089C | VPS21 | GTPase | 0.0067 | 1.0 | 0.968212 |
| YIR031C | DAL7 | malate synthase | 0.0068 | 1.0 | 0.952106 |
| YBR009C | HHF1 | DNA binding | 0.0078 | 1.0 | 0.952965 |
| YBR286W | APE3 | aminopeptidase | 0.0093 | 1.0 | 0.361603 |
| YLR155C | ASP3-1 | asparaginase | 0.0093 | 1.0 | 0.883431 |
| YCR009C | RVS161 | cytoskeletal protein binding | 0.0113 | 1.0 | 0.913926 |
| YDL232W | OST4 | protein binding | 0.0115 | 1.0 | 0.932106 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|--|----------------------------|-------------|-----------|
| YJL225C | | helicase | 0.0148 | 1.0 | 0.896168 |
| YKL056C | TMA19 | unknown | 0.0150 | 1.0 | 0.825276 |
| YFL006W | | | 0.0165 | 1.0 | 0.936657 |
| YBR120C | CBP6 | unknown | 0.0167 | 1.0 | 0.909303 |
| YLR448W | RPL6B | structural constituent of ribosome* | 0.0182 | 1.0 | 0.929721 |
| YJL071W | ARG2 | amino-acid N-acetyltransferase | 0.0197 | 1.0 | 0.901738 |
| YJL021C | | | 0.0200 | 1.0 | 0.764018 |
| YIL059C | | | 0.0215 | 1.0 | 0.752012 |
| YBR232C | | | 0.0217 | 1.0 | 0.907517 |
| YHR195W | NVJ1 | protein binding | 0.0220 | 1.0 | 0.849372 |
| YIR026C | YVH1 | protein tyrosine phosphatase * | 0.0225 | 1.0 | 0.953713 |
| YIL097W | FYV10 | unknown | 0.0227 | 1.0 | 0.797355 |
| YOL039W | RPP2A | structural constituent of ribosome | 0.0240 | 1.0 | 0.798216 |
| YBR125C | PTC4 | protein phosphatase type 2C | 0.0257 | 1.0 | 0.867112 |
| YDL097C | RPN6 | structural molecule | 0.0258 | 1.0 | 0.851374 |
| YHL021C | FMP12 | unknown | 0.0267 | 1.0 | 0.741442 |
| YIR008C | PRI1 | alpha DNA polymerase | 0.0277 | 1.0 | 0.874719 |
| YBR073W | RDH54 | DNA-dependent ATPase * | 0.0283 | 1.0 | 0.69516 |
| YNL031C | HHT2 | DNA binding | 0.0305 | 1.0 | 0.814157 |
| YIL143C | SSL2 | general RNA polymerase II transcription factor * | 0.0308 | 1.0 | 0.779992 |
| YLR375W | STP3 | unknown | 0.0333 | 1.0 | 0.571388 |
| YNL149C | PGA2 | unknown | 0.0335 | 1.0 | 0.782749 |
| YDL075W | RPL31A | structural constituent of ribosome | 0.0347 | 1.0 | 0.555455 |
| YGR098C | ESP1 | cysteine-type endopeptidase | 0.0348 | 1.0 | 0.925403 |
| YBR010W | HHT1 | DNA binding | 0.0360 | 1.0 | 0.863754 |
| YEL020C | | unknown | 0.0385 | 1.0 | 0.790251 |
| YBR100W | | | 0.0405 | 1.0 | 0.865627 |
| YER043C | SAH1 | adenosylhomocysteinase | 0.0430 | 1.0 | 0.676812 |
| YHR178W | STB5 | transcription factor | 0.0432 | 1.0 | 0.597239 |
| YHR198C | FMP22 | unknown | 0.0438 | 1.0 | 0.698314 |
| YDL081C | RPP1A | structural constituent of ribosome | 0.0443 | 1.0 | 0.894563 |
| YFR016C | | unknown | 0.0443 | 1.0 | 0.783366 |
| YAL054C | ACS1 | acetate-CoA ligase | 0.0445 | 1.0 | 0.853367 |
| YOL027C | MDM38 | unknown | 0.0470 | 1.0 | 0.580167 |
| YMR044W | IOC4 | protein binding | 0.0487 | 1.0 | 0.699442 |
| YNL197C | WHI3 | RNA binding | 0.0493 | 1.0 | 0.752765 |
| YLR157C | ASP3-2 | asparaginase | 0.0513 | 1.0 | 0.559126 |
| YGL221C | NIF3 | unknown | 0.0517 | 1.0 | 0.884325 |
| YIL177C | | helicase | 0.0523 | 1.0 | 0.712893 |
| YPR043W | RPL43A | structural constituent of ribosome | 0.0528 | 1.0 | 0.5684 |
| YOR053W | | | 0.0530 | 1.0 | 0.519903 |
| YIL146C | ECM37 | unknown | 0.0538 | 1.0 | 0.70498 |
| YBL058W | SHP1 | protein phosphatase type 1 regulator | 0.0540 | 1.0 | 0.652117 |
| YIL036W | CST6 | specific RNA polymerase II transcription factor | 0.0558 | 1.0 | 0.625373 |
| YDL234C | GYP7 | Rab GTPase activator | 0.0567 | 1.0 | 0.620932 |
| YJL176C | SWI3 | general RNA polymerase II transcription factor | 0.0570 | 1.0 | 0.149109 |
| YIL026C | IRR1 | protein binding | 0.0572 | 1.0 | 0.790645 |
| YJL149W | | unknown | 0.0572 | 1.0 | 0.784475 |
| YIL090W | ICE2 | unknown | 0.0573 | 1.0 | 0.815598 |
| YBR134W | | | 0.0583 | 1.0 | 0.717538 |
| YKL180W | RPL17A | structural constituent of ribosome | 0.0588 | 1.0 | 0.713634 |
| YCR107W | AAD3 | aryl-alcohol dehydrogenase | 0.0595 | 1.0 | 0.723654 |
| YOR315W | SFG1 | specific RNA polymerase II transcription factor | 0.0608 | 1.0 | 0.882602 |
| YBR079C | RPG1 | translation initiation factor | 0.0623 | 1.0 | 0.849129 |
| YPL235W | RVB2 | ATPase * | 0.0623 | 1.0 | 0.636872 |
| YLL019C | KNS1 | protein serine/threonine kinase | 0.0632 | 1.0 | 0.731721 |
| YNL115C | | unknown | 0.0638 | 1.0 | 0.400503 |
| YNL305C | | unknown | 0.0643 | 1.0 | 0.625113 |
| YBR203W | COS111 | unknown | 0.0650 | 1.0 | 0.830386 |
| YGR148C | RPL24B | structural constituent of ribosome* | 0.0650 | 1.0 | 0.592386 |
| YPR042C | PUF2 | mRNA binding | 0.0662 | 1.0 | 0.709883 |
| YPR148C | | unknown | 0.0662 | 1.0 | 0.197285 |
| YFL007W | BLM10 | proteasome activator | 0.0673 | 1.0 | 0.788506 |
| YOR285W | | unknown | 0.0678 | 1.0 | 0.564548 |
| YKL171W | | protein kinase | 0.0680 | 1.0 | 0.85326 |
| YCL050C | APA1 | bis(5'-nucleosyl)-tetraphosphatase | 0.0682 | 1.0 | 0.698987 |
| YAL053W | FLC2 | unknown | 0.0683 | 1.0 | 0.867804 |
| YJL110C | GZF3 | transcription factor * | 0.0688 | 1.0 | 0.559607 |
| YGL227W | VID30 | unknown | 0.0708 | 1.1 | 0.0128979 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YBR176W | ECM31 | 3-methyl-2-oxobutanoate hydroxymethyltransferase | 0.0718 | 1.1 | 0.239951 |
| YAR014C | BUD14 | protein phosphatase type 1 regulator | 0.0735 | 1.1 | 0.31085 |
| YFL021C-A | | | 0.0735 | 1.1 | 0.291533 |
| YKR002W | PAP1 | polynucleotide adenylyltransferase | 0.0742 | 1.1 | 0.83087 |
| YOR194C | TOA1 | general RNA polymerase II transcription factor | 0.0742 | 1.1 | 0.366512 |
| YCR051W | | unknown | 0.0767 | 1.1 | 0.691641 |
| YGL189C | RPS26A | structural constituent of ribosome | 0.0767 | 1.1 | 0.392322 |
| YNL016W | PUB1 | mRNA binding | 0.0792 | 1.1 | 0.523818 |
| YDR330W | UBX5 | unknown | 0.0808 | 1.1 | 0.481804 |
| YOL026C | MIM1 | unknown | 0.0813 | 1.1 | 0.241457 |
| YCL016C | DCC1 | unknown | 0.0815 | 1.1 | 0.429242 |
| YOR372C | NDD1 | transcriptional activator | 0.0820 | 1.1 | 0.510075 |
| YEL001C | | unknown | 0.0827 | 1.1 | 0.439822 |
| YJR074W | MOG1 | Ran GTPase binding | 0.0833 | 1.1 | 0.376654 |
| YPL260W | | unknown | 0.0850 | 1.1 | 0.533537 |
| YBR177C | EHT1 | serine hydrolase | 0.0873 | 1.1 | 0.206581 |
| YIL112W | HOS4 | NAD-dependent histone deacetylase * | 0.0878 | 1.1 | 0.336896 |
| YBL101C | ECM21 | unknown | 0.0880 | 1.1 | 0.130741 |
| YAL056W | GPB2 | signal transducer | 0.0887 | 1.1 | 0.581451 |
| YAL016W | TPD3 | protein phosphatase type 2A | 0.0898 | 1.1 | 0.558172 |
| YBR181C | RPS6B | structural constituent of ribosome | 0.0905 | 1.1 | 0.0788305 |
| YCL042W | | unknown | 0.0910 | 1.1 | 0.652731 |
| YPL219W | PCL8 | cyclin-dependent protein kinase | 0.0918 | 1.1 | 0.143769 |
| YGR117C | | unknown | 0.0930 | 1.1 | 0.482923 |
| YOR043W | WHI2 | phosphatase activator | 0.0932 | 1.1 | 0.563934 |
| YKR042W | UTH1 | unknown | 0.0937 | 1.1 | 0.695054 |
| YDR418W | RPL12B | structural constituent of ribosome | 0.0938 | 1.1 | 0.520595 |
| YBR175W | SWD3 | histone lysine N-methyltransferase (H3-K4 specific) | 0.0948 | 1.1 | 0.704378 |
| YGL058W | RAD6 | ubiquitin conjugating enzyme | 0.0953 | 1.1 | 0.729575 |
| YCR053W | THR4 | threonine synthase | 0.0988 | 1.1 | 0.370164 |
| YDL099W | BUG1 | unknown | 0.0992 | 1.1 | 0.263134 |
| YKL093W | MBR1 | unknown | 0.1000 | 1.1 | 0.43462 |
| YJL020C | BBC1 | myosin I binding | 0.1002 | 1.1 | 0.430835 |
| YMR152W | YIM1 | unknown | 0.1007 | 1.1 | 0.260185 |
| YIL107C | PFK26 | 6-phosphofructo-2-kinase | 0.1013 | 1.1 | 0.185519 |
| YER021W | RPN3 | unknown | 0.1018 | 1.1 | 0.180763 |
| YPL081W | RPS9A | structural constituent of ribosome | 0.1022 | 1.1 | 0.367343 |
| YMR282C | AEP2 | unknown | 0.1050 | 1.1 | 0.46097 |
| YCL003W | | | 0.1057 | 1.1 | 0.526202 |
| YNL042W | BOP3 | unknown | 0.1057 | 1.1 | 0.721034 |
| YPL222W | FMP40 | unknown | 0.1070 | 1.1 | 0.649179 |
| YLR237W | THI7 | thiamin transporter | 0.1075 | 1.1 | 0.737274 |
| YHR203C | RPS4B | structural constituent of ribosome | 0.1080 | 1.1 | 0.206443 |
| YEL060C | PRB1 | serine-type endopeptidase | 0.1083 | 1.1 | 0.081507 |
| YBR109C | CMD1 | protein binding | 0.1085 | 1.1 | 0.498652 |
| YJL191W | RPS14B | structural constituent of ribosome* | 0.1095 | 1.1 | 0.384347 |
| YDR233C | RTN1 | unknown | 0.1103 | 1.1 | 0.2583 |
| YJL140W | RPB4 | DNA-directed RNA polymerase | 0.1115 | 1.1 | 0.150697 |
| YDR263C | DIN7 | nuclease | 0.1117 | 1.1 | 0.142271 |
| YKL006W | RPL14A | structural constituent of ribosome* | 0.1145 | 1.1 | 0.165822 |
| YBR127C | VMA2 | hydrogen-transporting ATPase, rotational mechanism | 0.1152 | 1.1 | 0.609987 |
| YFL043C | | | 0.1157 | 1.1 | 0.45262 |
| YHR132C | ECM14 | metalloendopeptidase | 0.1158 | 1.1 | 0.364983 |
| YDL218W | | unknown | 0.1182 | 1.1 | 0.705368 |
| YLR178C | TFS1 | lipid binding | 0.1185 | 1.1 | 0.294337 |
| YBR131W | CCZ1 | guanyl-nucleotide exchange factor | 0.1193 | 1.1 | 0.252601 |
| YNL030W | HHF2 | DNA binding | 0.1203 | 1.1 | 0.325061 |
| YLR454W | FMP27 | unknown | 0.1212 | 1.1 | 0.307638 |
| YBL081W | | unknown | 0.1222 | 1.1 | 0.225264 |
| YNR047W | | protein kinase | 0.1223 | 1.1 | 0.643167 |
| YMR002W | | unknown | 0.1235 | 1.1 | 0.447 |
| YFR008W | FAR7 | unknown | 0.1240 | 1.1 | 0.272748 |
| YER160C | | | 0.1245 | 1.1 | 0.585023 |
| YER159C | BUR6 | transcription corepressor | 0.1247 | 1.1 | 0.0176098 |
| YBR260C | RGD1 | Rho GTPase activator | 0.1248 | 1.1 | 0.18176 |
| YDL140C | RPO21 | DNA-directed RNA polymerase | 0.1250 | 1.1 | 0.41749 |
| YPR016C | TIF6 | unknown | 0.1258 | 1.1 | 0.269512 |
| YOR251C | | thiosulfate sulfurtransferase | 0.1267 | 1.1 | 0.398732 |
| YCR042C | TAF2 | general RNA polymerase II transcription factor | 0.1268 | 1.1 | 0.318437 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|----------|-----------|---|----------------------------|-------------|-----------|
| YER080W | FMP29 | unknown | 0.1282 | 1.1 | 0.436598 |
| YCL049C | | unknown | 0.1292 | 1.1 | 0.290869 |
| YJR002W | MPP10 | unknown | 0.1295 | 1.1 | 0.348597 |
| YDR062W | LCB2 | serine C-palmitoyltransferase | 0.1300 | 1.1 | 0.478307 |
| YER100W | UBC6 | ubiquitin conjugating enzyme | 0.1328 | 1.1 | 0.138897 |
| YJL072C | PSF2 | unknown | 0.1348 | 1.1 | 0.398719 |
| YDR500C | RPL37B | structural constituent of ribosome | 0.1353 | 1.1 | 0.362215 |
| YMR168C | CEP3 | DNA bending * | 0.1358 | 1.1 | 0.183822 |
| YOR099W | KTR1 | alpha-1,2-mannosyltransferase | 0.1363 | 1.1 | 0.0384063 |
| YMR105C | PGM2 | phosphoglucosyltransferase | 0.1378 | 1.1 | 0.589437 |
| YCR059C | YIH1 | unknown | 0.1385 | 1.1 | 0.288575 |
| YDR217C | RAD9 | protein binding | 0.1387 | 1.1 | 0.0942463 |
| YDR309C | GIC2 | small GTPase regulator | 0.1408 | 1.1 | 0.189636 |
| YKR071C | DRE2 | unknown | 0.1410 | 1.1 | 0.61076 |
| YDR473C | PRP3 | RNA splicing factor , transesterification mechanism | 0.1412 | 1.1 | 0.295316 |
| YBR136W | MEC1 | protein kinase | 0.1452 | 1.1 | 0.549302 |
| YMR236W | TAF9 | general RNA polymerase II transcription factor | 0.1475 | 1.1 | 0.170939 |
| YER074W | RPS24A | structural constituent of ribosome | 0.1482 | 1.1 | 0.301609 |
| YJL015C | | | 0.1482 | 1.1 | 0.271545 |
| YMR143W | RPS16A | structural constituent of ribosome | 0.1513 | 1.1 | 0.405787 |
| YDR035W | ARO3 | 3-deoxy-7-phosphoheptulonate synthase | 0.1528 | 1.1 | 0.544787 |
| YNL118C | DCP2 | mRNA binding | 0.1532 | 1.1 | 0.46221 |
| YOR120W | GCY1 | aldo-keto reductase | 0.1545 | 1.1 | 0.29807 |
| YJL165C | HAL5 | protein kinase | 0.1572 | 1.1 | 0.223009 |
| YBR081C | SPT7 | structural molecule | 0.1573 | 1.1 | 0.210179 |
| YGL243W | TAD1 | tRNA specific adenosine deaminase | 0.1573 | 1.1 | 0.743083 |
| YHR039BC | | | 0.1575 | 1.1 | 0.270592 |
| YBR263W | SHM1 | glycine hydroxymethyltransferase | 0.1590 | 1.1 | 0.110675 |
| YAL019W | FUN30 | unknown | 0.1595 | 1.1 | 0.119214 |
| YDL045C | FAD1 | FMN adenyltransferase | 0.1602 | 1.1 | 0.28881 |
| YOR182C | RPS30B | structural constituent of ribosome | 0.1607 | 1.1 | 0.397334 |
| YBR249C | ARO4 | 3-deoxy-7-phosphoheptulonate synthase | 0.1613 | 1.1 | 0.300001 |
| YHR190W | ERG9 | farnesyl-diphosphate farnesyltransferase | 0.1613 | 1.1 | 0.0929202 |
| YBR215W | HPC2 | transcription regulator | 0.1620 | 1.1 | 0.159109 |
| YOR312C | RPL20B | structural constituent of ribosome | 0.1628 | 1.1 | 0.382632 |
| YBR290W | BSD2 | unknown | 0.1632 | 1.1 | 0.0225144 |
| YAL058W | CNE1 | unfolded protein binding | 0.1640 | 1.1 | 0.469466 |
| YKL063C | | unknown | 0.1657 | 1.1 | 0.178941 |
| YML045W | | | 0.1665 | 1.1 | 0.0178715 |
| YDR099W | BMH2 | protein binding | 0.1668 | 1.1 | 0.26254 |
| YBR212W | NGR1 | RNA binding | 0.1672 | 1.1 | 0.0951895 |
| YEL054C | RPL12A | structural constituent of ribosome | 0.1682 | 1.1 | 0.378921 |
| YDR516C | EMI2 | unknown | 0.1692 | 1.1 | 0.316558 |
| YOR377W | ATF1 | alcohol O-acetyltransferase | 0.1702 | 1.1 | 0.451847 |
| YDR130C | FIN1 | unknown | 0.1708 | 1.1 | 0.382842 |
| YCL002C | | unknown | 0.1713 | 1.1 | 0.165626 |
| YLR256W | HAP1 | specific RNA polymerase II transcription factor | 0.1713 | 1.1 | 0.397353 |
| YPR132W | RPS23B | structural constituent of ribosome | 0.1733 | 1.1 | 0.0253757 |
| YCR032W | BPH1 | unknown | 0.1745 | 1.1 | 0.291174 |
| YNL117W | MLS1 | malate synthase | 0.1755 | 1.1 | 0.148843 |
| YBR160W | CDC28 | cyclin-dependent protein kinase | 0.1757 | 1.1 | 0.108491 |
| YDL165W | CDC36 | 3'-5'-exoribonuclease | 0.1762 | 1.1 | 0.25734 |
| YER182W | FMP10 | unknown | 0.1762 | 1.1 | 0.476109 |
| YOL139C | CDC33 | translation initiation factor | 0.1768 | 1.1 | 0.269887 |
| YER049W | TPA1 | unknown | 0.1770 | 1.1 | 0.166726 |
| YAR042W | SWH1 | phosphatidylinositol binding | 0.1777 | 1.1 | 0.82101 |
| YOL011W | PLB3 | lysophospholipase | 0.1777 | 1.1 | 0.0345318 |
| YDL083C | RPS16B | structural constituent of ribosome | 0.1780 | 1.1 | 0.137987 |
| YKR066C | CCP1 | cytochrome-c peroxidase | 0.1790 | 1.1 | 0.200555 |
| YHR191C | CTF8 | unknown | 0.1793 | 1.1 | 0.0080886 |
| YIL121W | QDR2 | multidrug efflux pump | 0.1793 | 1.1 | 0.0350443 |
| YIL043C | CBR1 | cytochrome-b5 reductase | 0.1803 | 1.1 | 0.442247 |
| YAL021C | CCR4 | 3'-5'-exoribonuclease | 0.1822 | 1.1 | 0.380068 |
| YBL049W | MOH1 | unknown | 0.1840 | 1.1 | 0.199181 |
| YPL156C | PRM4 | unknown | 0.1840 | 1.1 | 0.38523 |
| YBR289W | SNF5 | general RNA polymerase II transcription factor | 0.1845 | 1.1 | 0.15961 |
| YKL122C | SRP21 | signal sequence binding | 0.1845 | 1.1 | 0.201753 |
| YML024W | RPS17A | structural constituent of ribosome | 0.1848 | 1.1 | 0.345004 |
| YER079W | | unknown | 0.1855 | 1.1 | 0.51985 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YDL131W | LYS21 | homocitrate synthase | 0.1857 | 1.1 | 0.132558 |
| YML026C | RPS18B | structural constituent of ribosome | 0.1867 | 1.1 | 0.124204 |
| YMR016C | SOK2 | transcription factor | 0.1867 | 1.1 | 0.157004 |
| YKL006C-A | SFT1 | v-SNARE | 0.1887 | 1.1 | 0.287021 |
| Q0060 | AI3 | endonuclease | 0.1905 | 1.1 | 0.0746155 |
| YDL084W | SUB2 | protein binding | 0.1907 | 1.1 | 0.0291847 |
| YBR295W | PCA1 | ATPase , coupled to transmembrane movement of ions, phosph | 0.1922 | 1.1 | 0.0819892 |
| YAR015W | ADE1 | phosphoribosylaminoimidazolesuccinocarboxamide synthase | 0.1925 | 1.1 | 0.008313 |
| YJL177W | RPL17B | structural constituent of ribosome | 0.1933 | 1.1 | 0.0645773 |
| YMR125W | STO1 | mRNA binding | 0.1945 | 1.1 | 0.187596 |
| YCR082W | AHC2 | unknown | 0.1947 | 1.1 | 0.327742 |
| YLR158C | ASP3-3 | asparaginase | 0.1962 | 1.1 | 0.224188 |
| YBL085W | BOI1 | phospholipid binding | 0.1968 | 1.1 | 0.179566 |
| YHL011C | PRS3 | ribose phosphate diphosphokinase | 0.1978 | 1.1 | 0.268141 |
| YDL016C | | | 0.2007 | 1.1 | 0.299784 |
| YOR293W | RPS10A | structural constituent of ribosome | 0.2055 | 1.2 | 0.13611 |
| YBL037W | APL3 | unknown | 0.2075 | 1.2 | 0.130578 |
| YAL017W | PSK1 | protein serine/threonine kinase | 0.2080 | 1.2 | 0.0869401 |
| YGR046W | | unknown | 0.2082 | 1.2 | 0.123359 |
| YBR119W | MUD1 | RNA binding | 0.2092 | 1.2 | 0.130848 |
| YML042W | CAT2 | carnitine O-acetyltransferase | 0.2097 | 1.2 | 0.0453306 |
| YLR168C | | unknown | 0.2102 | 1.2 | 0.0088148 |
| YCR036W | RBK1 | ATP binding | 0.2103 | 1.2 | 0.112399 |
| YDL085W | NDE2 | NADH dehydrogenase | 0.2115 | 1.2 | 0.0451916 |
| YLR438W | CAR2 | ornithine-oxo-acid transaminase | 0.2117 | 1.2 | 0.213545 |
| YLR441C | RPS1A | structural constituent of ribosome | 0.2117 | 1.2 | 0.175398 |
| YNL078W | NIS1 | unknown | 0.2125 | 1.2 | 0.213784 |
| YMR116C | ASC1 | unknown | 0.2133 | 1.2 | 0.167254 |
| YCR002C | CDC10 | GTPase * | 0.2155 | 1.2 | 0.0145124 |
| YER019C-A | SBH2 | protein transporter | 0.2155 | 1.2 | 0.394379 |
| YJR007W | SUI2 | translation initiation factor | 0.2155 | 1.2 | 0.47037 |
| YOR321W | PMT3 | dolichyl-phosphate-mannose-protein mannosyltransferase | 0.2162 | 1.2 | 0.227859 |
| YBR048W | RPS11B | structural constituent of ribosome | 0.2163 | 1.2 | 0.0904646 |
| YOR317W | FAA1 | long-chain-fatty-acid-CoA ligase | 0.2165 | 1.2 | 0.0842238 |
| YOR064C | YNG1 | histone acetyltransferase | 0.2182 | 1.2 | 0.0350022 |
| YPL131W | RPL5 | structural constituent of ribosome* | 0.2188 | 1.2 | 0.174037 |
| YBR066C | NRG2 | transcriptional repressor | 0.2190 | 1.2 | 0.161822 |
| YPL220W | RPL1A | structural constituent of ribosome | 0.2193 | 1.2 | 0.227748 |
| YER020W | GPA2 | GTPase | 0.2198 | 1.2 | 0.0893473 |
| YER019W | ISC1 | phospholipase C | 0.2207 | 1.2 | 0.325469 |
| YBR287W | ZSP1 | unknown | 0.2227 | 1.2 | 0.0494068 |
| YER081W | SER3 | phosphoglycerate dehydrogenase | 0.2242 | 1.2 | 0.0045936 |
| YBR051W | | | 0.2257 | 1.2 | 0.0517947 |
| YJR100C | | unknown | 0.2273 | 1.2 | 0.414574 |
| YMR064W | AEP1 | unknown | 0.2275 | 1.2 | 0.0462257 |
| YJL127C | SPT10 | histone acetyltransferase | 0.2283 | 1.2 | 0.498844 |
| YBR253W | SRB6 | RNA polymerase II transcription mediator | 0.2293 | 1.2 | 0.278533 |
| YHL033C | RPL8A | structural constituent of ribosome | 0.2300 | 1.2 | 0.137696 |
| YCR057C | PWP2 | snoRNA binding | 0.2303 | 1.2 | 0.0348982 |
| YDL010W | | unknown | 0.2318 | 1.2 | 0.272822 |
| YKR098C | UBP11 | ubiquitin-specific protease | 0.2318 | 1.2 | 0.0463327 |
| YLR327C | TMA10 | unknown | 0.2330 | 1.2 | 0.1726 |
| YGL237C | HAP2 | transcriptional activator | 0.2335 | 1.2 | 0.0793536 |
| YKL032C | IXR1 | DNA binding | 0.2340 | 1.2 | 0.315165 |
| YBR169C | SSE2 | unknown | 0.2353 | 1.2 | 0.0641767 |
| YAL009W | SPO7 | unknown | 0.2362 | 1.2 | 0.565516 |
| YLR238W | FAR10 | unknown | 0.2372 | 1.2 | 0.106673 |
| YDR511W | ACN9 | unknown | 0.2378 | 1.2 | 0.252092 |
| YMR212C | EFR3 | unknown | 0.2383 | 1.2 | 0.652341 |
| YIL077C | | unknown | 0.2407 | 1.2 | 0.226894 |
| YCR031 | | | 0.2413 | 1.2 | 0.046791 |
| YMR051C | | | 0.2425 | 1.2 | 0.099358 |
| YHR006W | STP2 | specific RNA polymerase II transcription factor | 0.2432 | 1.2 | 0.133915 |
| YDR266C | | unknown | 0.2435 | 1.2 | 0.128454 |
| YFL-TYB | | | 0.2442 | 1.2 | 0.103945 |
| YLR264W | RPS28B | structural constituent of ribosome | 0.2450 | 1.2 | 0.202117 |
| YCR045C | | serine-type peptidase | 0.2452 | 1.2 | 0.577389 |
| YGL122C | NAB2 | poly(A) binding | 0.2478 | 1.2 | 0.170935 |
| YDR077W | SED1 | structural constituent of cell wall | 0.2483 | 1.2 | 0.294513 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YDL226C | GCS1 | actin binding | 0.2485 | 1.2 | 0.0310436 |
| YDR068W | DOS2 | unknown | 0.2495 | 1.2 | 0.0837498 |
| YDR071C | PAA1 | aralkylamine N-acetyltransferase * | 0.2505 | 1.2 | 0.0933319 |
| YPR086W | SUA7 | general RNA polymerase II transcription factor | 0.2505 | 1.2 | 0.113247 |
| YBR049C | REB1 | RNA polymerase II transcription factor * | 0.2507 | 1.2 | 0.092407 |
| YDL223C | HBT1 | unknown | 0.2525 | 1.2 | 0.0393533 |
| YIL021W | RPB3 | DNA-directed RNA polymerase | 0.2527 | 1.2 | 0.0074357 |
| YLR206W | ENT2 | clathrin binding | 0.2537 | 1.2 | 0.0017845 |
| YDR092W | UBC13 | ubiquitin conjugating enzyme | 0.2542 | 1.2 | 0.309298 |
| YPL079W | RPL21B | structural constituent of ribosome | 0.2547 | 1.2 | 0.195736 |
| YMR110C | HFD1 | 3-chloroallyl aldehyde dehydrogenase | 0.2548 | 1.2 | 0.0151451 |
| YDR090C | | unknown | 0.2555 | 1.2 | 0.0702783 |
| YFL016C | MDJ1 | unfolded protein binding | 0.2562 | 1.2 | 0.0070729 |
| YBR071W | | unknown | 0.2563 | 1.2 | 0.270707 |
| YMR046C | | | 0.2570 | 1.2 | 0.13098 |
| YDL186W | | unknown | 0.2580 | 1.2 | 0.0272931 |
| YDR324C | UTP4 | snoRNA binding | 0.2580 | 1.2 | 0.124495 |
| YNL023C | FAP1 | transcription factor | 0.2583 | 1.2 | 0.716553 |
| YAL033W | POP5 | ribonuclease P * | 0.2590 | 1.2 | 0.0767786 |
| YDR167W | TAF10 | general RNA polymerase II transcription factor | 0.2598 | 1.2 | 0.0630406 |
| YCR008W | SAT4 | protein kinase | 0.2605 | 1.2 | 0.0558661 |
| YCR024C-A | PMP1 | enzyme regulator | 0.2607 | 1.2 | 0.0841958 |
| YIR010W | DSN1 | unknown | 0.2610 | 1.2 | 0.085467 |
| YDL005C | MED2 | RNA polymerase II transcription mediator | 0.2617 | 1.2 | 0.177827 |
| YCR044C | PER1 | unknown | 0.2620 | 1.2 | 0.521016 |
| YIL135C | VHS2 | unknown | 0.2637 | 1.2 | 0.2734 |
| YBR157C | ICS2 | unknown | 0.2648 | 1.2 | 0.0172777 |
| YJR076C | CDC11 | structural constituent of cytoskeleton* | 0.2657 | 1.2 | 0.197752 |
| YDL092W | SRP14 | signal sequence binding | 0.2658 | 1.2 | 0.0113611 |
| YBR115C | LYS2 | L-aminoadipate-semialdehyde dehydrogenase | 0.2660 | 1.2 | 0.502537 |
| YER014W | HEM14 | protoporphyrinogen oxidase | 0.2673 | 1.2 | 0.0026705 |
| YHR213W | | unknown | 0.2693 | 1.2 | 0.0410299 |
| YDR088C | SLU7 | RNA splicing factor , transesterification mechanism* | 0.2710 | 1.2 | 0.144377 |
| YNL154C | YCK2 | casein kinase I | 0.2715 | 1.2 | 0.0243373 |
| YDR533C | HSP31 | unfolded protein binding | 0.2738 | 1.2 | 0.0956208 |
| YLR325C | RPL38 | structural constituent of ribosome | 0.2748 | 1.2 | 0.304616 |
| YKL150W | MCR1 | cytochrome-b5 reductase | 0.2753 | 1.2 | 0.0699198 |
| YLR094C | GIS3 | unknown | 0.2768 | 1.2 | 0.253976 |
| YBR031W | RPL4A | structural constituent of ribosome | 0.2827 | 1.2 | 0.119102 |
| YLR040C | | unknown | 0.2835 | 1.2 | 0.0655853 |
| YLR250W | SSP120 | unknown | 0.2835 | 1.2 | 0.150086 |
| YCR062W | | | 0.2858 | 1.2 | 0.0008532 |
| YDR356W | SPC110 | structural constituent of cytoskeleton | 0.2862 | 1.2 | 0.370205 |
| YFR029W | PTR3 | amino acid binding | 0.2868 | 1.2 | 0.0125929 |
| YLR246W | ERF2 | protein-cysteine S-palmitoleyltransferase | 0.2898 | 1.2 | 0.0354252 |
| YDL091C | UBX3 | unknown | 0.2912 | 1.2 | 0.18028 |
| YDL190C | UFD2 | ubiquitin conjugating enzyme | 0.2912 | 1.2 | 0.0933201 |
| YGR146C | | unknown | 0.2915 | 1.2 | 0.0253532 |
| YLR093C | NYV1 | v-SNARE | 0.2920 | 1.2 | 0.120326 |
| YBR123C | TFC1 | RNA polymerase III transcription factor | 0.2928 | 1.2 | 0.168292 |
| YJR058C | APS2 | unknown | 0.2942 | 1.2 | 0.0314166 |
| YBR256C | RIB5 | riboflavin synthase | 0.2980 | 1.2 | 0.912071 |
| YJR123W | RPS5 | structural constituent of ribosome | 0.2980 | 1.2 | 0.199123 |
| YOL106W | | | 0.2987 | 1.2 | 0.0515506 |
| YDR012W | RPL4B | structural constituent of ribosome | 0.2990 | 1.2 | 0.0087733 |
| YCR025C | | | 0.3017 | 1.2 | 0.601868 |
| YDL106C | PHO2 | transcription factor | 0.3022 | 1.2 | 0.274582 |
| YKL172W | EBP2 | unknown | 0.3027 | 1.2 | 0.0001707 |
| YOR113W | AZF1 | DNA binding | 0.3033 | 1.2 | 0.0429644 |
| YOR184W | SER1 | phosphoserine transaminase | 0.3038 | 1.2 | 0.252962 |
| YPL206C | | glycerophosphodiester phosphodiesterase | 0.3067 | 1.2 | 0.0830177 |
| YGL123W | RPS2 | structural constituent of ribosome | 0.3077 | 1.2 | 0.0147807 |
| YLL045C | RPL8B | structural constituent of ribosome | 0.3085 | 1.2 | 0.0219904 |
| YMR194W | RPL36A | structural constituent of ribosome* | 0.3085 | 1.2 | 0.0337608 |
| YBR166C | TYR1 | prephenate dehydrogenase | 0.3087 | 1.2 | 0.384681 |
| YBR190W | | | 0.3098 | 1.2 | 0.0853908 |
| YJL069C | UTP18 | unknown | 0.3103 | 1.2 | 0.28073 |
| YGR007W | MUQ1 | ethanolamine-phosphate cytidyltransferase | 0.3117 | 1.2 | 0.0214033 |
| YJL164C | TPK1 | protein serine/threonine kinase | 0.3117 | 1.2 | 0.0565868 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YMR250W | GAD1 | glutamate decarboxylase | 0.3117 | 1.2 | 0.320592 |
| YCL066W | HMLALPHA1 | transcription coactivator | 0.3118 | 1.2 | 0.0915848 |
| YDR269C | | | 0.3132 | 1.2 | 0.0544512 |
| YBR113W | | | 0.3140 | 1.2 | 0.0322234 |
| YGL095C | VPS45 | unfolded protein binding | 0.3142 | 1.2 | 0.107025 |
| YDL188C | PPH22 | protein phosphatase type 2A | 0.3148 | 1.2 | 0.0332659 |
| YDL020C | RPN4 | transcriptional activator | 0.3157 | 1.2 | 0.0720395 |
| YDL048C | STP4 | unknown | 0.3163 | 1.2 | 0.0312737 |
| YBR112C | CYC8 | transcription coactivator * | 0.3167 | 1.2 | 0.0276492 |
| YDR350C | TCM10 | unknown | 0.3178 | 1.2 | 0.0509807 |
| YNL162W | RPL42A | structural constituent of ribosome | 0.3182 | 1.2 | 0.0600358 |
| YHR149C | SKG6 | unknown | 0.3195 | 1.2 | 0.0068105 |
| YBR168W | PEX32 | unknown | 0.3200 | 1.2 | 0.279415 |
| YGL126W | SCS3 | unknown | 0.3210 | 1.2 | 0.369375 |
| YDR180W | SCC2 | unknown | 0.3217 | 1.2 | 0.222937 |
| YCL041C | | | 0.3228 | 1.3 | 0.0152257 |
| YNL166C | BNI5 | unknown | 0.3257 | 1.3 | 0.0390715 |
| YDL161W | ENT1 | clathrin binding | 0.3258 | 1.3 | 0.0197532 |
| YOR095C | RKI1 | ribose-5-phosphate isomerase | 0.3270 | 1.3 | 0.0710363 |
| YMR235C | RNA1 | Ran GTPase activator | 0.3272 | 1.3 | 0.0349585 |
| YMR022W | QRI8 | ubiquitin conjugating enzyme | 0.3298 | 1.3 | 0.0583425 |
| YBR012W-A | | | 0.3308 | 1.3 | 0.116522 |
| YBR084C-A | RPL19A | structural constituent of ribosome | 0.3312 | 1.3 | 0.0315089 |
| YDR055W | PST1 | unknown | 0.3312 | 1.3 | 0.0099697 |
| YIR006C | PAN1 | protein binding | 0.3312 | 1.3 | 0.1055 |
| YJR003C | | unknown | 0.3320 | 1.3 | 0.110934 |
| YGR229C | SMI1 | unknown | 0.3332 | 1.3 | 0.392184 |
| YBR172C | SMY2 | unknown | 0.3352 | 1.3 | 0.0035667 |
| Q0140 | VAR1 | structural constituent of ribosome | 0.3353 | 1.3 | 0.0080991 |
| YOR122C | PFY1 | actin monomer binding | 0.3353 | 1.3 | 0.0569877 |
| YNL161W | CBK1 | protein kinase | 0.3357 | 1.3 | 0.220263 |
| YHR141C | RPL42B | structural constituent of ribosome | 0.3362 | 1.3 | 0.104475 |
| YBL041W | PRE7 | endopeptidase | 0.3365 | 1.3 | 0.142208 |
| YOR389W | | unknown | 0.3390 | 1.3 | 0.286057 |
| YAL060W | BDH1 | (R,R)-butanediol dehydrogenase | 0.3405 | 1.3 | 0.0019043 |
| YLR197W | SIK1 | unknown | 0.3407 | 1.3 | 0.104173 |
| YBR089W | | | 0.3420 | 1.3 | 0.0224113 |
| YJL076W | NET1 | rDNA binding | 0.3433 | 1.3 | 0.107821 |
| YNR002C | ATO2 | ammonium transporter | 0.3453 | 1.3 | 0.0120224 |
| YML040W | | | 0.3457 | 1.3 | 0.0432481 |
| YIL074C | SER33 | phosphoglycerate dehydrogenase | 0.3465 | 1.3 | 0.0189035 |
| YDR153C | ENT5 | clathrin binding | 0.3468 | 1.3 | 0.0712946 |
| YGR256W | GND2 | phosphogluconate dehydrogenase (decarboxylating) | 0.3468 | 1.3 | 0.255532 |
| YBR229C | ROT2 | alpha-glucosidase | 0.3470 | 1.3 | 0.292602 |
| YHR010W | RPL27A | structural constituent of ribosome | 0.3478 | 1.3 | 0.138529 |
| YGR019W | UGA1 | 4-aminobutyrate transaminase | 0.3487 | 1.3 | 0.138266 |
| YLR309C | JMH1 | unknown | 0.3500 | 1.3 | 0.031425 |
| YOR304W | ISW2 | ATPase | 0.3502 | 1.3 | 0.473588 |
| YMR097C | MTG1 | GTPase | 0.3505 | 1.3 | 0.0806746 |
| YLR301W | | unknown | 0.3507 | 1.3 | 0.0976603 |
| YNR057C | BIO4 | dethiobiotin synthase | 0.3512 | 1.3 | 0.0454883 |
| YIL149C | MLP2 | ribonucleoprotein binding | 0.3520 | 1.3 | 0.0097889 |
| YDR476C | | unknown | 0.3537 | 1.3 | 0.0146229 |
| YDR067C | OCA6 | unknown | 0.3550 | 1.3 | 0.0281261 |
| YOR311C | HSD1 | unknown | 0.3553 | 1.3 | 0.234589 |
| YOR128C | ADE2 | phosphoribosylaminoimidazole carboxylase | 0.3563 | 1.3 | 0.0312847 |
| YER102W | RPS8B | structural constituent of ribosome | 0.3567 | 1.3 | 0.0143663 |
| YJL147C | | unknown | 0.3570 | 1.3 | 0.218333 |
| YGR251W | | unknown | 0.3577 | 1.3 | 0.0348247 |
| YDR060W | MAK21 | unknown | 0.3580 | 1.3 | 0.0141598 |
| YER109C | FLO8 | specific RNA polymerase II transcription factor | 0.3585 | 1.3 | 0.0026077 |
| YGR147C | NAT2 | peptide alpha-N-acetyltransferase | 0.3587 | 1.3 | 0.0049887 |
| YPL239W | YAR1 | unknown | 0.3588 | 1.3 | 0.0012312 |
| YOR096W | RPS7A | structural constituent of ribosome | 0.3603 | 1.3 | 0.045111 |
| YCL034W | LSB5 | unknown | 0.3610 | 1.3 | 0.172697 |
| YBR189W | RPS9B | structural constituent of ribosome | 0.3612 | 1.3 | 0.0013001 |
| YML074C | FPR3 | peptidyl-prolyl cis-trans isomerase | 0.3613 | 1.3 | 0.0369408 |
| YOL082W | ATG19 | protein binding | 0.3625 | 1.3 | 0.140094 |
| YJR066W | TOR1 | protein binding | 0.3627 | 1.3 | 0.0301496 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|---|----------------------------|-------------|-----------|
| YDR298C | ATP5 | structural molecule * | 0.3642 | 1.3 | 0.0285107 |
| YCR061W(1) | | | 0.3643 | 1.3 | 0.181657 |
| YDL243C | AAD4 | aryl-alcohol dehydrogenase | 0.3650 | 1.3 | 0.167374 |
| YPL061W | ALD6 | 3-chloroallyl aldehyde dehydrogenase | 0.3653 | 1.3 | 0.0460235 |
| YER068W | MOT2 | 3'-5'-exoribonuclease * | 0.3665 | 1.3 | 0.0115801 |
| YIL048W | NEO1 | phospholipid-translocating ATPase * | 0.3673 | 1.3 | 0.105522 |
| YDR250C | | | 0.3683 | 1.3 | 0.0733077 |
| YGL103W | RPL28 | structural constituent of ribosome* | 0.3708 | 1.3 | 0.114567 |
| YHR146W | CRP1 | DNA binding | 0.3727 | 1.3 | 0.0301264 |
| YNR052C | POP2 | 3'-5'-exoribonuclease | 0.3727 | 1.3 | 0.0213755 |
| YOR359W | VTS1 | RNA binding | 0.3737 | 1.3 | 0.075993 |
| YMR315W | unknown | | 0.3740 | 1.3 | 0.138719 |
| YGR014W | MSB2 | osmosensor | 0.3742 | 1.3 | 0.568245 |
| YBR297W | MAL33 | transcription factor | 0.3743 | 1.3 | 0.040974 |
| YBR264C | YPT10 | GTPase * | 0.3757 | 1.3 | 0.0980819 |
| Q0255 | unknown | | 0.3760 | 1.3 | 0.0575876 |
| YDR452W | PPN1 | endopolyphosphatase | 0.3777 | 1.3 | 0.0742538 |
| YBR230C | OM14 | unknown | 0.3818 | 1.3 | 0.139045 |
| YCR052W | RSC6 | unknown | 0.3820 | 1.3 | 0.111431 |
| YCR047C | BUD23 | S-adenosylmethionine-dependent methyltransferase | 0.3822 | 1.3 | 0.166106 |
| YLL010C | PSR1 | phosphoprotein phosphatase | 0.3825 | 1.3 | 0.0219582 |
| YDL124W | | alpha-keto amide reductase * | 0.3832 | 1.3 | 0.0152227 |
| YMR272C | SCS7 | oxidoreductase | 0.3838 | 1.3 | 0.079297 |
| YMR163C | | unknown | 0.3840 | 1.3 | 0.191744 |
| YDL166C | FAP7 | nucleoside-triphosphatase | 0.3845 | 1.3 | 0.16134 |
| YPR036W | VMA13 | hydrogen-transporting ATPase , rotational mechanism | 0.3852 | 1.3 | 0.0018528 |
| YHR202W | | unknown | 0.3853 | 1.3 | 0.0792575 |
| YOR298W | MUM3 | acyltransferase | 0.3858 | 1.3 | 0.364013 |
| YDL072C | YET3 | unknown | 0.3872 | 1.3 | 0.0332545 |
| YLR177W | | unknown | 0.3898 | 1.3 | 0.015294 |
| YCL011C | GBP2 | RNA binding | 0.3902 | 1.3 | 0.085295 |
| YGR027C | RPS25A | structural constituent of ribosome | 0.3923 | 1.3 | 0.0417248 |
| YOR276W | CAF20 | translation regulator | 0.3940 | 1.3 | 0.0374735 |
| YML051W | GAL80 | transcription corepressor | 0.3943 | 1.3 | 0.0583907 |
| YJR095W | SFC1 | succinate:fumarate antiporter | 0.3953 | 1.3 | 0.113921 |
| YDL121C | | unknown | 0.3958 | 1.3 | 0.209981 |
| YLR104W | | unknown | 0.3968 | 1.3 | 0.0283164 |
| YPL199C | | unknown | 0.3980 | 1.3 | 0.0073836 |
| YIL113W | SDP1 | MAP kinase | 0.3990 | 1.3 | 0.0187789 |
| YHL001W | RPL14B | structural constituent of ribosome* | 0.3993 | 1.3 | 0.0243855 |
| YCL024W | KCC4 | protein kinase | 0.4015 | 1.3 | 0.164817 |
| YLR384C | IK3 | RNA polymerase II transcription elongation factor * | 0.4022 | 1.3 | 0.235114 |
| YMR070W | MOT3 | DNA binding | 0.4023 | 1.3 | 0.0828388 |
| YDL061C | RPS29B | structural constituent of ribosome | 0.4025 | 1.3 | 0.226787 |
| YDL125C | HNT1 | hydrolase * | 0.4062 | 1.3 | 0.110569 |
| YPL052W | OAZ1 | enzyme inhibitor | 0.4068 | 1.3 | 0.0534972 |
| YJL017W | | | 0.4070 | 1.3 | 0.118868 |
| YPL273W | SAM4 | homocysteine S-methyltransferase | 0.4070 | 1.3 | 0.489671 |
| YER056C-A | RPL34A | structural constituent of ribosome | 0.4083 | 1.3 | 0.0114097 |
| YCR029C | | | 0.4095 | 1.3 | 0.197745 |
| YLR333C | RPS25B | structural constituent of ribosome | 0.4108 | 1.3 | 0.164936 |
| YOR167C | RPS28A | structural constituent of ribosome | 0.4112 | 1.3 | 0.0826127 |
| YDL142C | CRD1 | cardiolipin synthase | 0.4140 | 1.3 | 0.0718081 |
| YOL040C | RPS15 | structural constituent of ribosome | 0.4155 | 1.3 | 0.117769 |
| YDL217C | TIM22 | protein transporter | 0.4157 | 1.3 | 0.0730385 |
| YMR242C | RPL20A | structural constituent of ribosome | 0.4162 | 1.3 | 0.0949676 |
| YDL225W | SHS1 | structural constituent of cytoskeleton | 0.4175 | 1.3 | 0.0136115 |
| YER027C | GAL83 | AMP-activated protein kinase | 0.4177 | 1.3 | 0.0304636 |
| YML113W | DAT1 | AT DNA binding | 0.4195 | 1.3 | 0.0660129 |
| YBR129C | OPY1 | unknown | 0.4197 | 1.3 | 0.0607726 |
| YKL186C | MTR2 | protein binding | 0.4205 | 1.3 | 0.0382333 |
| YDR072C | IPT1 | transferase , transferring phosphorus-containing groups | 0.4208 | 1.3 | 0.0306595 |
| YIL104C | SHQ1 | unknown | 0.4217 | 1.3 | 0.053 |
| YDR351W | SBE2 | unknown | 0.4218 | 1.3 | 0.0174488 |
| YGL080W | FMP37 | unknown | 0.4225 | 1.3 | 0.0517235 |
| YLR335W | NUP2 | structural molecule | 0.4232 | 1.3 | 0.0215867 |
| YBR062C | | unknown | 0.4258 | 1.3 | 0.0737451 |
| YFL-TYA | | | 0.4278 | 1.3 | 0.139567 |
| YMR142C | RPL13B | structural constituent of ribosome | 0.4288 | 1.3 | 0.026996 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YDL046W | NPC2 | unknown | 0.4290 | 1.3 | 0.0765605 |
| YPL143W | RPL33A | structural constituent of ribosome | 0.4312 | 1.3 | 0.0870198 |
| YER088C | DOT6 | unknown | 0.4340 | 1.4 | 0.0029541 |
| YBR214W | SDS24 | unknown | 0.4342 | 1.4 | 0.0276292 |
| YGR009C | SEC9 | t-SNARE | 0.4355 | 1.4 | 0.12963 |
| YBR114W | RAD16 | DNA-dependent ATPase | 0.4360 | 1.4 | 0.0626064 |
| YLR383W | SMC6 | unknown | 0.4365 | 1.4 | 0.135411 |
| YOR173W | DCS2 | unknown | 0.4365 | 1.4 | 0.0686687 |
| YIL027C | KRE27 | unknown | 0.4372 | 1.4 | 0.107721 |
| YBR106W | PHO88 | phosphate transporter | 0.4373 | 1.4 | 0.0943904 |
| YHR134W | WSS1 | unknown | 0.4375 | 1.4 | 0.0212561 |
| YHR209W | | S-adenosylmethionine-dependent methyltransferase | 0.4387 | 1.4 | 0.0411227 |
| YBR086C | IST2 | unknown | 0.4388 | 1.4 | 0.076393 |
| YLR052W | IES3 | unknown | 0.4407 | 1.4 | 0.0376626 |
| YCLX10C | | | 0.4412 | 1.4 | 0.0877594 |
| YNL021W | HDA1 | histone deacetylase | 0.4420 | 1.4 | 0.185184 |
| YFL058W | THI5 | unknown | 0.4427 | 1.4 | 0.0851235 |
| YJL030W | MAD2 | unknown | 0.4427 | 1.4 | 0.0465429 |
| YKR091W | SRL3 | unknown | 0.4467 | 1.4 | 0.0030785 |
| YOL126C | MDH2 | L-malate dehydrogenase | 0.4482 | 1.4 | 0.0095309 |
| YOR283W | | unknown | 0.4502 | 1.4 | 0.10103 |
| YHR163W | SOL3 | 6-phosphogluconolactonase | 0.4517 | 1.4 | 0.0070757 |
| YJR012C | | unknown | 0.4520 | 1.4 | 0.0161885 |
| YBR130C | SHE3 | mRNA binding | 0.4523 | 1.4 | 0.0319372 |
| YOR245C | DGA1 | diacylglycerol O-acyltransferase | 0.4527 | 1.4 | 0.0843416 |
| YJL117W | PHO86 | unknown | 0.4548 | 1.4 | 0.0008273 |
| YBL092W | RPL32 | structural constituent of ribosome | 0.4552 | 1.4 | 0.258129 |
| YDR447C | RPS17B | structural constituent of ribosome | 0.4555 | 1.4 | 0.0391831 |
| YJL064W | | | 0.4560 | 1.4 | 0.121646 |
| YLR053C | | unknown | 0.4563 | 1.4 | 0.0201412 |
| YHR030C | SLT2 | MAP kinase | 0.4572 | 1.4 | 0.102423 |
| YBR167C | POP7 | ribonuclease P * | 0.4577 | 1.4 | 0.0125835 |
| YLR253W | | unknown | 0.4580 | 1.4 | 0.0403765 |
| YDR145W | TAF12 | general RNA polymerase II transcription factor | 0.4590 | 1.4 | 0.0394653 |
| YDR022C | CIS1 | unknown | 0.4603 | 1.4 | 0.496707 |
| YDR513W | GRX2 | thiol-disulfide exchange intermediate * | 0.4603 | 1.4 | 0.0796488 |
| TDR261C-D | | | 0.4617 | 1.4 | 0.131645 |
| YMR073C | | unknown | 0.4617 | 1.4 | 0.198626 |
| YJL141C | YAK1 | protein kinase | 0.4620 | 1.4 | 0.0264559 |
| YBR128C | ATG14 | unknown | 0.4643 | 1.4 | 0.0699366 |
| YHR152W | SPO12 | unknown | 0.4655 | 1.4 | 0.0200978 |
| YBR075W | | | 0.4662 | 1.4 | 0.0065598 |
| YLR410W-B | | | 0.4668 | 1.4 | 0.0081619 |
| YOR046C | DBP5 | RNA helicase | 0.4673 | 1.4 | 0.11785 |
| YBR191W | RPL21A | structural constituent of ribosome | 0.4682 | 1.4 | 0.0187838 |
| YGL077C | HNMI | choline transporter | 0.4682 | 1.4 | 0.0295363 |
| YBR138C | | unknown | 0.4685 | 1.4 | 0.223255 |
| YDL088C | ASM4 | structural molecule | 0.4685 | 1.4 | 0.0179667 |
| YAR037W | | | 0.4697 | 1.4 | 0.246423 |
| YBR213W | MET8 | ferrochelatase | 0.4697 | 1.4 | 0.012594 |
| YLR382C | NAM2 | mRNA binding | 0.4697 | 1.4 | 0.0349826 |
| YDR510W | SMT3 | protein tag | 0.4700 | 1.4 | 0.0711014 |
| YIR023W | DAL81 | specific RNA polymerase II transcription factor | 0.4707 | 1.4 | 0.0145459 |
| YDL082W | RPL13A | structural constituent of ribosome | 0.4708 | 1.4 | 0.0195166 |
| YKL028W | TFA1 | general RNA polymerase II transcription factor | 0.4708 | 1.4 | 0.0946728 |
| YIL116W | HIS5 | histidinol-phosphate transaminase | 0.4710 | 1.4 | 0.0323842 |
| YDR034C | LYS14 | transcriptional activator | 0.4713 | 1.4 | 0.088733 |
| YIL045W | PIG2 | protein phosphatase type 1 regulator | 0.4730 | 1.4 | 0.0305551 |
| YML053C | | unknown | 0.4745 | 1.4 | 0.0084153 |
| YIL133C | RPL16A | structural constituent of ribosome* | 0.4758 | 1.4 | 0.0315509 |
| YDL177C | | unknown | 0.4765 | 1.4 | 0.0381995 |
| YIL111W | COX5B | cytochrome-c oxidase | 0.4767 | 1.4 | 0.262548 |
| YHR135C | YCK1 | casein kinase | 0.4797 | 1.4 | 0.0532657 |
| YDL053C | PBP4 | unknown | 0.4837 | 1.4 | 0.0087411 |
| YPL095C | EEB1 | unknown | 0.4838 | 1.4 | 0.0188507 |
| YDR471W | RPL27B | structural constituent of ribosome | 0.4853 | 1.4 | 0.0868826 |
| YOL133W | HRT1 | protein binding | 0.4855 | 1.4 | 0.130335 |
| YOR145C | PNO1 | unfolded protein binding | 0.4863 | 1.4 | 0.139019 |
| YLR388W | RPS29A | structural constituent of ribosome | 0.4890 | 1.4 | 0.07198 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YOR054C | VHS3 | phosphopantothenoylcysteine decarboxylase * | 0.4910 | 1.4 | 0.027021 |
| YOL068C | HST1 | NAD-dependent histone deacetylase * | 0.4915 | 1.4 | 0.0738916 |
| YMR243C | ZRC1 | di-, tri-valent inorganic cation transporter | 0.4918 | 1.4 | 0.0656337 |
| YFL067W | | unknown | 0.4932 | 1.4 | 0.64905 |
| YBL101W-A | | | 0.4947 | 1.4 | 0.004212 |
| YLR330W | CHS5 | unknown | 0.4953 | 1.4 | 0.0196172 |
| YDR450W | RPS18A | structural constituent of ribosome | 0.4958 | 1.4 | 0.0149866 |
| YER036C | ARB1 | ATPase | 0.4958 | 1.4 | 0.0280296 |
| YBL040C | ERD2 | HDEL sequence binding | 0.4973 | 1.4 | 0.028403 |
| YGL135W | RPL1B | structural constituent of ribosome | 0.4973 | 1.4 | 0.0214487 |
| YDL079C | MRK1 | glycogen synthase kinase | 0.4982 | 1.4 | 0.0958719 |
| YPR041W | TIF5 | translation initiation factor * | 0.4995 | 1.4 | 0.101006 |
| YER065C | ICL1 | isocitrate lyase | 0.4997 | 1.4 | 0.0095397 |
| YCR041W | | | 0.4998 | 1.4 | 0.2451 |
| YIL105C | SLM1 | phosphoinositide binding | 0.5017 | 1.4 | 0.0071928 |
| YJL118W | | unknown | 0.5033 | 1.4 | 0.0547805 |
| YNL178W | RPS3 | structural constituent of ribosome | 0.5055 | 1.4 | 0.0043822 |
| YDR025W | RPS11A | structural constituent of ribosome | 0.5063 | 1.4 | 0.0552956 |
| YML058W | SML1 | enzyme inhibitor | 0.5070 | 1.4 | 0.12005 |
| YBL097W | BRN1 | unknown | 0.5075 | 1.4 | 0.281202 |
| YIL069C | RPS24B | structural constituent of ribosome | 0.5078 | 1.4 | 0.007055 |
| YKL156W | RPS27A | structural constituent of ribosome | 0.5095 | 1.4 | 0.0562405 |
| YBR140C | IRA1 | Ras GTPase activator | 0.5097 | 1.4 | 0.0292406 |
| YIL007C | NAS2 | unknown | 0.5100 | 1.4 | 0.0793274 |
| YAR010C | | | 0.5145 | 1.4 | 0.0581187 |
| YBL072C | RPS8A | structural constituent of ribosome | 0.5162 | 1.4 | 0.189202 |
| YDL025C | | protein kinase | 0.5205 | 1.4 | 0.146723 |
| YIR012W | SQT1 | unknown | 0.5222 | 1.4 | 0.0949048 |
| YBL064C | PRX1 | thioredoxin peroxidase | 0.5225 | 1.4 | 0.0425174 |
| YOL098C | | metalloendopeptidase | 0.5232 | 1.4 | 0.33902 |
| YJR028W | | | 0.5252 | 1.4 | 0.0315859 |
| YGL128C | CWC23 | unknown | 0.5255 | 1.4 | 0.0029366 |
| YDL191W | RPL35A | structural constituent of ribosome | 0.5270 | 1.4 | 0.0553661 |
| YCR084C | TUP1 | general transcriptional repressor | 0.5277 | 1.4 | 0.017961 |
| YDR408C | ADE8 | phosphoribosylglycinamide formyltransferase | 0.5307 | 1.4 | 0.0435428 |
| YJL065C | DLS1 | unknown | 0.5308 | 1.4 | 0.018932 |
| YNL027W | CRZ1 | transcription factor | 0.5322 | 1.4 | 0.0157153 |
| YLL023C | | unknown | 0.5328 | 1.4 | 0.037853 |
| YHR097C | | unknown | 0.5340 | 1.4 | 0.036475 |
| YBR135W | CKS1 | protein kinase | 0.5357 | 1.4 | 0.0125355 |
| YAR002W | NUP60 | structural constituent of nuclear pore | 0.5368 | 1.5 | 0.0145382 |
| YFL031W | HAC1 | transcription factor * | 0.5368 | 1.5 | 0.181046 |
| YDR508C | GNP1 | amino acid transporter | 0.5378 | 1.5 | 0.0344047 |
| YBL005W-A | | | 0.5390 | 1.5 | 0.0549268 |
| YMR124W | | unknown | 0.5398 | 1.5 | 0.0144626 |
| YDR354W | TRP4 | anthranilate phosphoribosyltransferase | 0.5402 | 1.5 | 0.0161396 |
| YNL244C | SUI1 | translation initiation factor | 0.5410 | 1.5 | 0.0709532 |
| YNL099C | OCA1 | protein tyrosine phosphatase | 0.5412 | 1.5 | 0.0227138 |
| YER148W | SPT15 | DNA binding | 0.5420 | 1.5 | 0.0299641 |
| YMR197C | VTI1 | v-SNARE | 0.5445 | 1.5 | 0.0362647 |
| YNL059C | ARP5 | unknown | 0.5455 | 1.5 | 0.061866 |
| YLR290C | | unknown | 0.5472 | 1.5 | 0.0954273 |
| YPL049C | DIG1 | DNA binding | 0.5475 | 1.5 | 0.237246 |
| YCR038C | BUD5 | signal transducer * | 0.5492 | 1.5 | 0.0310974 |
| YHR122W | | unknown | 0.5495 | 1.5 | 0.0016139 |
| YDL224C | WHI4 | RNA binding | 0.5498 | 1.5 | 0.0645596 |
| YHL002W | HSE1 | protein binding | 0.5508 | 1.5 | 0.0751863 |
| YHR154W | RTT107 | unknown | 0.5520 | 1.5 | 0.050038 |
| YLR198C | | | 0.5523 | 1.5 | 0.0246025 |
| YKL065C | YET1 | unknown | 0.5527 | 1.5 | 0.0526849 |
| YNL133C | FYV6 | unknown | 0.5528 | 1.5 | 0.0089595 |
| YOR052C | | unknown | 0.5535 | 1.5 | 0.0426082 |
| YBR050C | REG2 | protein phosphatase type 1 regulator | 0.5555 | 1.5 | 0.0287962 |
| YBR030W | | unknown | 0.5583 | 1.5 | 0.112549 |
| YDL047W | SIT4 | protein serine/threonine phosphatase | 0.5598 | 1.5 | 0.0045282 |
| YAR027W | UIP3 | unknown | 0.5613 | 1.5 | 0.0202892 |
| YDL134C | PPH21 | protein phosphatase type 2A | 0.5613 | 1.5 | 0.025438 |
| YLR029C | RPL15A | structural constituent of ribosome* | 0.5618 | 1.5 | 0.0170283 |
| YFL059W | SNZ3 | protein binding | 0.5623 | 1.5 | 0.0841028 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|------------|-----------|--|----------------------------|-------------|-----------|
| YDR154C | | | 0.5638 | 1.5 | 0.0412276 |
| YBR024W | SCO2 | thioredoxin peroxidase | 0.5688 | 1.5 | 0.0025094 |
| YDR492W | IZH1 | metal ion binding | 0.5695 | 1.5 | 0.0337674 |
| YPR040W | TIP41 | unknown | 0.5732 | 1.5 | 0.0498624 |
| YDR412W | RRP17 | unknown | 0.5737 | 1.5 | 0.0097174 |
| YJL189W | RPL39 | structural constituent of ribosome | 0.5743 | 1.5 | 0.0179627 |
| YIL004C | BET1 | v-SNARE | 0.5752 | 1.5 | 0.0578862 |
| YDR244W | PEX5 | peroxisome targeting sequence binding | 0.5757 | 1.5 | 0.013908 |
| YBR279W | PAF1 | RNA polymerase II transcription elongation factor | 0.5770 | 1.5 | 0.0056696 |
| YKL012W | PRP40 | RNA binding | 0.5770 | 1.5 | 0.0371327 |
| YDL134C-A | | | 0.5793 | 1.5 | 0.0077575 |
| YDL169C | UGX2 | unknown | 0.5795 | 1.5 | 0.011808 |
| YLL039C | UBI4 | protein tag* | 0.5797 | 1.5 | 0.0067012 |
| YJL190C | RPS22A | structural constituent of ribosome | 0.5798 | 1.5 | 0.0301632 |
| YDR121W | DPB4 | epsilon DNA polymerase | 0.5802 | 1.5 | 0.00634 |
| YBR211C | AME1 | unknown | 0.5813 | 1.5 | 0.141025 |
| YJR096W | | aldo-keto reductase * | 0.5823 | 1.5 | 0.0362997 |
| YLR391W | | | 0.5835 | 1.5 | 0.0889035 |
| YCL005W | LDB16 | unknown | 0.5843 | 1.5 | 0.124221 |
| YER026C | CHO1 | CDP-diacylglycerol-serine O-phosphatidyltransferase | 0.5845 | 1.5 | 0.002478 |
| YDR064W | RPS13 | structural constituent of ribosome | 0.5860 | 1.5 | 0.0312294 |
| YHR142W | CHS7 | unknown | 0.5893 | 1.5 | 0.0109231 |
| YAR009C | | | 0.5893 | 1.5 | 0.0463007 |
| YOR103C | OST2 | dolichyl-diphosphooligosaccharide-protein glycotransferase | 0.5898 | 1.5 | 0.0175108 |
| YKL214C | YRA2 | RNA binding | 0.5900 | 1.5 | 0.642384 |
| YCR020C-A | MAK31 | peptide alpha-N-acetyltransferase | 0.5912 | 1.5 | 0.0017026 |
| YGL078C | DBP3 | ATP-dependent RNA helicase | 0.5917 | 1.5 | 0.0085144 |
| YDL136W | RPL35B | structural constituent of ribosome | 0.5922 | 1.5 | 0.0330865 |
| YMR255W | GFD1 | unknown | 0.6005 | 1.5 | 0.01597 |
| YPL054W | LEE1 | unknown | 0.6010 | 1.5 | 0.0335987 |
| YLR075W | RPL10 | structural constituent of ribosome | 0.6032 | 1.5 | 0.0106968 |
| YDL070W | BDF2 | unknown | 0.6053 | 1.5 | 0.0078577 |
| YPR102C | RPL11A | structural constituent of ribosome | 0.6055 | 1.5 | 0.0026455 |
| YOR234C | RPL33B | structural constituent of ribosome | 0.6063 | 1.5 | 0.0912172 |
| YNL096C | RPS7B | structural constituent of ribosome | 0.6070 | 1.5 | 0.0665578 |
| YDR432W | NPL3 | mRNA binding | 0.6080 | 1.5 | 0.0993888 |
| YPL237W | SUI3 | translation initiation factor | 0.6088 | 1.5 | 0.0963182 |
| YBR162W-A | YSY6 | unknown | 0.6102 | 1.5 | 0.015102 |
| YBR077C | SLM4 | phosphatidylinositol-3,4-bisphosphate binding | 0.6107 | 1.5 | 0.0166044 |
| YOL149W | DCP1 | mRNA binding | 0.6113 | 1.5 | 0.0041306 |
| YDR498C | SEC20 | v-SNARE | 0.6113 | 1.5 | 0.0097686 |
| YAL014C | SYN8 | SNAP receptor | 0.6118 | 1.5 | 0.0402033 |
| YHR214C-B | | | 0.6175 | 1.5 | 0.0107341 |
| YDL077C | VAM6 | Rab guanyl-nucleotide exchange factor | 0.6188 | 1.5 | 0.0043489 |
| YMR005W | TAF4 | RNA polymerase II transcription factor | 0.6192 | 1.5 | 0.002214 |
| YNL062C | GCD10 | tRNA binding | 0.6198 | 1.5 | 0.0047471 |
| YJL136C | RPS21B | structural constituent of ribosome | 0.6203 | 1.5 | 0.0064227 |
| YJL018W | | | 0.6252 | 1.5 | 0.0458914 |
| YLR332W | MID2 | transmembrane receptor | 0.6278 | 1.5 | 0.0162083 |
| YAL031C | GIP4 | unknown | 0.6280 | 1.5 | 0.0331935 |
| YBL036C | | alanine racemase | 0.6283 | 1.5 | 0.0350416 |
| YPL229W | | unknown | 0.6293 | 1.5 | 0.0434246 |
| YDR343C | HXT6 | glucose transporter | 0.6313 | 1.5 | 0.0460598 |
| YAL015C | NTG1 | DNA-(apurinic or apyrimidinic site) lyase | 0.6333 | 1.6 | 0.0641866 |
| YPL238C | | | 0.6363 | 1.6 | 0.0291676 |
| YLL027W | ISA1 | unknown | 0.6368 | 1.6 | 0.0044931 |
| YDR433W | | | 0.6375 | 1.6 | 0.0037363 |
| YNL116W | DMA2 | unknown | 0.6427 | 1.6 | 0.0118315 |
| YOL073C | | unknown | 0.6428 | 1.6 | 0.0332842 |
| YIL040W | APQ12 | unknown | 0.6450 | 1.6 | 0.0455902 |
| YCR069W(1) | | | 0.6453 | 1.6 | 0.0242917 |
| YBL032W | HEK2 | mRNA binding | 0.6462 | 1.6 | 0.110691 |
| YGR137W | | | 0.6492 | 1.6 | 0.0098867 |
| YBL033C | RIB1 | cyclohydrolase | 0.6495 | 1.6 | 0.0430691 |
| YDL228C | | | 0.6510 | 1.6 | 0.0614825 |
| YCL039W | GID7 | unknown | 0.6512 | 1.6 | 0.0227334 |
| YDR069C | DOA4 | endopeptidase | 0.6525 | 1.6 | 0.0039496 |
| YOR152C | | unknown | 0.6605 | 1.6 | 0.101074 |
| YAR028W | | unknown | 0.6613 | 1.6 | 0.0181914 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YIL142C | | | 0.6645 | 1.6 | 0.0343115 |
| YDR373W | FRQ1 | enzyme activator | 0.6650 | 1.6 | 0.0351373 |
| YGR085C | RPL11B | structural constituent of ribosome | 0.6678 | 1.6 | 0.0083789 |
| YLR243W | | signal sequence binding | 0.6678 | 1.6 | 0.033278 |
| YOR244W | ESA1 | histone acetyltransferase | 0.6718 | 1.6 | 0.0142352 |
| YNL180C | RHO5 | GTPase | 0.6752 | 1.6 | 0.0137025 |
| YDR251W | PAM1 | unknown | 0.6755 | 1.6 | 0.360361 |
| YOR357C | SNX3 | protein binding | 0.6782 | 1.6 | 0.0322226 |
| YML097C | VPS9 | guanyl-nucleotide exchange factor | 0.6788 | 1.6 | 0.022509 |
| YCR070W | | | 0.6837 | 1.6 | 0.0191098 |
| YHR021C | RPS27B | structural constituent of ribosome | 0.6843 | 1.6 | 0.0513548 |
| YDL170W | UGA3 | transcription factor | 0.6923 | 1.6 | 0.0040976 |
| YCL044C | MGR1 | unknown | 0.6935 | 1.6 | 0.002867 |
| YHR150W | PEX28 | unknown | 0.6945 | 1.6 | 0.0076472 |
| YMR298W | LIP1 | sphingosine N-acyltransferase | 0.7003 | 1.6 | 0.10402 |
| YCRX16C | | | 0.7005 | 1.6 | 0.000299 |
| YMR191W | SPG5 | unknown | 0.7018 | 1.6 | 0.0222206 |
| YCR020C | PET18 | unknown | 0.7027 | 1.6 | 0.0132927 |
| YPR149W | NCE102 | unknown | 0.7047 | 1.6 | 0.0335362 |
| YMR096W | SNZ1 | protein binding | 0.7097 | 1.6 | 0.0041331 |
| YJR086W | STE18 | GTPase | 0.7127 | 1.6 | 0.0380325 |
| YLR254C | NDL1 | unknown | 0.7133 | 1.6 | 0.0487382 |
| YDL233W | | unknown | 0.7197 | 1.6 | 0.0115499 |
| YBR183W | YPC1 | ceramidase | 0.7220 | 1.6 | 0.0264319 |
| YGL166W | CUP2 | ligand-regulated transcription factor | 0.7227 | 1.7 | 0.0006974 |
| YNL001W | DOM34 | unknown | 0.7243 | 1.7 | 0.0397806 |
| YJL133W | MRS3 | iron ion transporter | 0.7258 | 1.7 | 0.03809 |
| YPR065W | ROX1 | specific transcriptional repressor | 0.7258 | 1.7 | 0.0016087 |
| YER013W | PRP22 | RNA splicing factor , transesterification mechanism | 0.7295 | 1.7 | 0.044427 |
| YIL148W | RPL40A | structural constituent of ribosome* | 0.7297 | 1.7 | 0.0006812 |
| YOL144W | NOP8 | unknown | 0.7337 | 1.7 | 0.0195138 |
| YDR073W | SNF11 | general RNA polymerase II transcription factor | 0.7363 | 1.7 | 0.0505516 |
| YPL004C | LSP1 | protein kinase | 0.7373 | 1.7 | 0.0104057 |
| YMR121C | RPL15B | structural constituent of ribosome | 0.7375 | 1.7 | 0.0032652 |
| YDL212W | SHR3 | unfolded protein binding | 0.7397 | 1.7 | 0.0051169 |
| YDL110C | TMA17 | unknown | 0.7413 | 1.7 | 0.0109625 |
| YLR076C | | | 0.7455 | 1.7 | 0.0111283 |
| YIL152W | | unknown | 0.7462 | 1.7 | 0.0106222 |
| YCR096C | HMRA2 | unknown | 0.7497 | 1.7 | 0.0059965 |
| YCR039C | MATALPHA2 | transcription corepressor | 0.7568 | 1.7 | 0.0099801 |
| YBR201W | DER1 | unknown | 0.7575 | 1.7 | 0.0230302 |
| YLR270W | DCS1 | hydrolase , acting on acid anhydrides, in phosphorus-containing | 0.7578 | 1.7 | 0.0234946 |
| YBR083W | TEC1 | specific RNA polymerase II transcription factor | 0.7617 | 1.7 | 0.0287579 |
| YOR385W | | unknown | 0.7630 | 1.7 | 0.0432233 |
| YHR170W | NMD3 | protein binding | 0.7645 | 1.7 | 0.0076408 |
| YGR248W | SOL4 | 6-phosphogluconolactonase | 0.7660 | 1.7 | 0.0018052 |
| YOR367W | SCP1 | protein binding | 0.7678 | 1.7 | 0.0269859 |
| YDR397C | NCB2 | transcription corepressor | 0.7748 | 1.7 | 0.0073126 |
| YCL067C | HMLALPHA2 | transcription corepressor | 0.7758 | 1.7 | 0.0176063 |
| YNL195C | | unknown | 0.7763 | 1.7 | 0.0208827 |
| YCL040W | GLK1 | glucokinase | 0.7798 | 1.7 | 0.0129569 |
| YIR017C | MET28 | DNA binding | 0.7800 | 1.7 | 0.0758299 |
| YIL052C | RPL34B | structural constituent of ribosome | 0.7853 | 1.7 | 0.0072553 |
| YEL074W | | | 0.7868 | 1.7 | 0.0377551 |
| YDR002W | YRB1 | Ran GTPase binding | 0.7923 | 1.7 | 0.0774179 |
| YNL156C | NSG2 | unknown | 0.7937 | 1.7 | 0.0455285 |
| YCR080W | | | 0.7945 | 1.7 | 0.0009706 |
| YDL182W | LYS20 | homocitrate synthase | 0.8002 | 1.7 | 0.0029764 |
| YHL015W | RPS20 | structural constituent of ribosome | 0.8023 | 1.7 | 0.0371291 |
| YBR182C | SMP1 | transcription factor | 0.8027 | 1.7 | 0.0142551 |
| YDL208W | NHP2 | RNA binding | 0.8077 | 1.8 | 0.0117274 |
| YPL057C | SUR1 | mannosyltransferase | 0.8095 | 1.8 | 0.0126662 |
| YIL060W | | unknown | 0.8175 | 1.8 | 0.0265058 |
| YHR145C | | | 0.8182 | 1.8 | 0.0196212 |
| YNL192W | CHS1 | chitin synthase | 0.8285 | 1.8 | 0.0439633 |
| YDR366C | | unknown | 0.8287 | 1.8 | 0.0163353 |
| YMR153W | NUP53 | structural molecule | 0.8300 | 1.8 | 0.0820494 |
| YGL070C | RPB9 | DNA-directed RNA polymerase | 0.8307 | 1.8 | 0.0511236 |
| YDL230W | PTP1 | protein tyrosine phosphatase | 0.8313 | 1.8 | 0.0067083 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YER087C-A | | | 0.8340 | 1.8 | 0.0864664 |
| YMR251W-A | HOR7 | unknown | 0.8340 | 1.8 | 0.0571777 |
| YCL031C | RRP7 | unknown | 0.8360 | 1.8 | 0.0141313 |
| YAR031W | PRM9 | unknown | 0.8370 | 1.8 | 0.0107993 |
| YIR016W | | unknown | 0.8392 | 1.8 | 0.0091005 |
| YCR005C | CIT2 | citrate (Si)-synthase | 0.8480 | 1.8 | 0.0523574 |
| YFR013W | IOC3 | protein binding | 0.8507 | 1.8 | 0.0511829 |
| YHR140W | | unknown | 0.8512 | 1.8 | 0.0062437 |
| YMR195W | ICY1 | unknown | 0.8540 | 1.8 | 0.0242994 |
| YNL119W | NCS2 | unknown | 0.8573 | 1.8 | 0.0219324 |
| YGR051C | | | 0.8588 | 1.8 | 0.132115 |
| YMR149W | SWP1 | dolichyl-diphosphooligosaccharide-protein glycotransferase | 0.8595 | 1.8 | 0.0499717 |
| YDL128W | VCX1 | calcium ion transporter | 0.8645 | 1.8 | 0.0129911 |
| YOR309C | | | 0.8673 | 1.8 | 0.0597032 |
| YDL227C | HO | endonuclease | 0.8693 | 1.8 | 0.0036788 |
| YOR091W | TMA46 | unknown | 0.8700 | 1.8 | 0.0095877 |
| YHR199C | FMP34 | unknown | 0.8758 | 1.8 | 0.0050944 |
| YDL129W | | unknown | 0.8773 | 1.8 | 0.0567164 |
| YOR348C | PUT4 | L-proline permease | 0.8805 | 1.8 | 0.0015679 |
| YDR306C | | protein binding | 0.8842 | 1.8 | 0.0438353 |
| YDL051W | LHP1 | RNA binding | 0.8850 | 1.8 | 0.0192443 |
| YPR113W | PIS1 | CDP-diacylglycerol-inositol 3-phosphatidyltransferase | 0.8887 | 1.9 | 0.0167012 |
| YOR104W | PIN2 | unknown | 0.8907 | 1.9 | 0.0079078 |
| YBR014C | | unknown | 0.8920 | 1.9 | 0.0201327 |
| YDL178W | SDH4 | succinate dehydrogenase (ubiquinone) | 0.8943 | 1.9 | 0.0296624 |
| YNL157W | IGO1 | unknown | 0.8953 | 1.9 | 0.0047717 |
| YHR049W | FSH1 | serine hydrolase | 0.8962 | 1.9 | 0.0206594 |
| YDR120C | TRM1 | tRNA (guanine-N2-)-methyltransferase | 0.8963 | 1.9 | 0.0147429 |
| YKL024C | URA6 | uridine kinase | 0.8965 | 1.9 | 0.0146188 |
| YDR203W | | | 0.9038 | 1.9 | 0.0259082 |
| YBR065C | ECM2 | unknown | 0.9042 | 1.9 | 0.0022371 |
| YJL192C | SOP4 | unknown | 0.9068 | 1.9 | 0.0025798 |
| YPL060W | LPE10 | magnesium ion transporter | 0.9075 | 1.9 | 0.0154464 |
| YHR155W | YSP1 | unknown | 0.9083 | 1.9 | 0.0443863 |
| YOR057W | SGT1 | chaperone binding | 0.9155 | 1.9 | 0.0253564 |
| YGR241C | YAP1802 | clathrin binding | 0.9222 | 1.9 | 0.0173609 |
| YGL172W | NUP49 | structural molecule | 0.9258 | 1.9 | 0.0074146 |
| YDR083W | RRP8 | methyltransferase | 0.9353 | 1.9 | 0.0271996 |
| YCR050C | | | 0.9420 | 1.9 | 0.0577847 |
| YOR316C | COT1 | di-, tri-valent inorganic cation transporter | 0.9422 | 1.9 | 0.0329205 |
| YJL089W | SIP4 | specific RNA polymerase II transcription factor | 0.9442 | 1.9 | 0.0099058 |
| YOR021C | | unknown | 0.9527 | 1.9 | 0.0186045 |
| YNR046W | TRM112 | zinc ion binding | 0.9548 | 1.9 | 0.0388793 |
| YLR107W | REX3 | 3'-5' exonuclease | 0.9560 | 1.9 | 0.0159249 |
| YOL109W | ZEO1 | unknown | 0.9563 | 1.9 | 0.0379235 |
| YIL144W | TID3 | structural constituent of cytoskeleton | 0.9583 | 1.9 | 0.0064321 |
| YNL191W | | unknown | 0.9645 | 2.0 | 0.0176501 |
| YGR136W | LSB1 | unknown | 0.9688 | 2.0 | 0.0222979 |
| YML128C | MSC1 | unknown | 0.9702 | 2.0 | 0.0105665 |
| YLR109W | AHP1 | thioredoxin peroxidase | 0.9712 | 2.0 | 0.0038796 |
| YDR074W | TPS2 | trehalose-phosphatase | 0.9730 | 2.0 | 0.0289385 |
| YGR034W | RPL26B | structural constituent of ribosome | 0.9808 | 2.0 | 0.0885126 |
| YLR022C | SDO1 | unknown | 0.9823 | 2.0 | 0.0091679 |
| YGL117W | | unknown | 0.9872 | 2.0 | 0.0133172 |
| YLR328W | NMA1 | nicotinamide-nucleotide adenyllyltransferase | 0.9883 | 2.0 | 0.0133009 |
| YHR187W | IKI1 | RNA polymerase II transcription elongation factor | 0.9890 | 2.0 | 0.007731 |
| YLR292C | SEC72 | protein transporter | 0.9930 | 2.0 | 0.0019322 |
| YBR034C | HMT1 | protein-arginine N-methyltransferase | 0.9942 | 2.0 | 0.0053895 |
| YKR025W | RPC37 | DNA-directed RNA polymerase | 0.9970 | 2.0 | 0.0389254 |
| YJL125C | GCD14 | tRNA (adenine-N1-)-methyltransferase | 1.0000 | 2.0 | 0.003202 |
| YHR043C | DOG2 | 2-deoxyglucose-6-phosphatase | 1.0052 | 2.0 | 0.0246612 |
| YGR230W | BNS1 | unknown | 1.0062 | 2.0 | 0.0216224 |
| YGR069W | | | 1.0148 | 2.0 | 0.0008504 |
| YJL060W | BNA3 | arylformamidase | 1.0352 | 2.0 | 0.0506385 |
| YKR040C | | | 1.0392 | 2.1 | 0.0097195 |
| YER025W | GCD11 | translation initiation factor | 1.0402 | 2.1 | 0.0365487 |
| YCR028C | FEN2 | pantothenate transporter | 1.0405 | 2.1 | 0.0236263 |
| YIL106W | MOB1 | kinase | 1.0433 | 2.1 | 0.0001591 |
| YLR252W | | | 1.0562 | 2.1 | 0.0404698 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YHR133C | NSG1 | protein binding | 1.0582 | 2.1 | 0.0038236 |
| YIL063C | YRB2 | unknown | 1.0637 | 2.1 | 0.0160592 |
| YDR256C | CTA1 | catalase | 1.0642 | 2.1 | 0.0422466 |
| YBR155W | CNS1 | unfolded protein binding | 1.0655 | 2.1 | 0.0022346 |
| YDL031W | DBP10 | ATP-dependent RNA helicase | 1.0657 | 2.1 | 0.0409663 |
| YCL065W | | | 1.0702 | 2.1 | 0.0234966 |
| YLR267W | BOP2 | unknown | 1.0703 | 2.1 | 0.0027182 |
| YCR033W | SNT1 | NAD-dependent histone deacetylase | 1.0733 | 2.1 | 0.0120961 |
| YDL222C | FMP45 | unknown | 1.0790 | 2.1 | 0.0101303 |
| YDL076C | RXT3 | unknown | 1.0815 | 2.1 | 0.0062744 |
| YDR527W | RBA50 | DNA-directed RNA polymerase | 1.0913 | 2.1 | 0.0138646 |
| YDL074C | BRE1 | ubiquitin-protein ligase | 1.0925 | 2.1 | 0.0071606 |
| YBR021W | FUR4 | uracil permease | 1.0927 | 2.1 | 0.0223852 |
| YDL123W | SNA4 | unknown | 1.0927 | 2.1 | 0.000194 |
| YAL059W | ECM1 | unknown | 1.1005 | 2.1 | 0.0392358 |
| YMR170C | ALD2 | 3-chloroallyl aldehyde dehydrogenase | 1.1058 | 2.2 | 0.0154439 |
| YHR171W | ATG7 | APG12 activating enzyme | 1.1080 | 2.2 | 0.0170534 |
| YPR025C | CCL1 | general RNA polymerase II transcription factor | 1.1100 | 2.2 | 0.109907 |
| YKL050C | | unknown | 1.1175 | 2.2 | 0.120823 |
| YNL194C | | unknown | 1.1220 | 2.2 | 0.0005191 |
| YMR262W | | unknown | 1.1272 | 2.2 | 0.0229439 |
| YKL151C | | unknown | 1.1275 | 2.2 | 0.0571066 |
| YBR165W | UBS1 | unknown | 1.1475 | 2.2 | 0.0023997 |
| YDR181C | SAS4 | acetyltransferase * | 1.1503 | 2.2 | 0.015197 |
| YOR023C | AHC1 | histone acetyltransferase | 1.1563 | 2.2 | 0.0270164 |
| YJL122W | ALB1 | unknown | 1.1597 | 2.2 | 0.0716043 |
| YHR138C | | endopeptidase inhibitor | 1.1625 | 2.2 | 0.0018937 |
| YNL112W | DBP2 | RNA helicase | 1.1717 | 2.3 | 0.0030629 |
| YJL144W | | unknown | 1.1767 | 2.3 | 0.0026914 |
| YCR035C | RRP43 | 3'-5'-exoribonuclease | 1.1787 | 2.3 | 0.0128518 |
| YCLX01W | | | 1.1955 | 2.3 | 0.011882 |
| YDR399W | HPT1 | hypoxanthine phosphoribosyltransferase | 1.1975 | 2.3 | 0.0464471 |
| YER024W | YAT2 | carnitine O-acetyltransferase | 1.2010 | 2.3 | 0.0089968 |
| YBL043W | ECM13 | unknown | 1.2050 | 2.3 | 0.0036862 |
| YIR018W | YAP5 | RNA polymerase II transcription factor | 1.2080 | 2.3 | 0.0159803 |
| YBL093C | ROX3 | RNA polymerase II transcription mediator | 1.2083 | 2.3 | 0.0245215 |
| YDR361C | BCP1 | unknown | 1.2132 | 2.3 | 0.0208792 |
| YNL143C | | | 1.2200 | 2.3 | 0.0046535 |
| YOR262W | | unknown | 1.2250 | 2.3 | 0.0104374 |
| YHR054C | | unknown | 1.2358 | 2.4 | 0.0104088 |
| YDR252W | BTT1 | unfolded protein binding | 1.2360 | 2.4 | 0.0011034 |
| YHR071W | PCL5 | cyclin-dependent protein kinase | 1.2392 | 2.4 | 0.0036633 |
| YLR251W | SYM1 | unknown | 1.2575 | 2.4 | 0.0037094 |
| YDL133W | | unknown | 1.2590 | 2.4 | 0.0009272 |
| YBR063C | | unknown | 1.2793 | 2.4 | 0.0501857 |
| YCR027C | RHB1 | GTPase | 1.2892 | 2.4 | 0.0481794 |
| YOR220W | WSP1 | unknown | 1.2978 | 2.5 | 0.102601 |
| YOL077C | BRX1 | rRNA primary transcript binding | 1.3110 | 2.5 | 0.0045937 |
| YGR057C | LST7 | protein transporter | 1.3122 | 2.5 | 0.0108821 |
| YDR037W | KRS1 | lysine-tRNA ligase | 1.3157 | 2.5 | 4.485E-05 |
| YCR016W | | unknown | 1.3220 | 2.5 | 0.0040942 |
| YLR414C | | unknown | 1.3288 | 2.5 | 0.0057038 |
| YEL026W | SNU13 | RNA binding | 1.3315 | 2.5 | 0.0466166 |
| YAL012W | CYS3 | cystathionine gamma-lyase | 1.3347 | 2.5 | 0.0485253 |
| YNL046W | | unknown | 1.3482 | 2.5 | 0.166658 |
| YHR161C | YAP1801 | clathrin binding | 1.3502 | 2.5 | 0.0003253 |
| YHR095W | | | 1.3560 | 2.6 | 0.0596932 |
| YLR061W | RPL22A | structural constituent of ribosome | 1.3692 | 2.6 | 0.0045889 |
| YNL002C | RLP7 | rRNA binding | 1.3708 | 2.6 | 0.007672 |
| YOL053C-A | | | 1.3720 | 2.6 | 0.014656 |
| YDR368W | YPR1 | oxidoreductase | 1.3863 | 2.6 | 0.0029524 |
| YBR105C | VID24 | unknown | 1.4198 | 2.7 | 0.0019282 |
| YPL135W | ISU1 | protein binding | 1.4242 | 2.7 | 0.0011447 |
| YHR104W | GRE3 | aldo-keto reductase * | 1.4313 | 2.7 | 0.0036065 |
| YML070W | DAK1 | glycerone kinase | 1.4385 | 2.7 | 0.0010459 |
| YDL204W | RTN2 | unknown | 1.4515 | 2.7 | 0.0033247 |
| YGR043C | | transaldolase | 1.4565 | 2.7 | 0.005118 |
| YGR086C | PIL1 | protein kinase | 1.4580 | 2.7 | 0.0033372 |
| YER150W | SPI1 | unknown | 1.4653 | 2.8 | 0.0089646 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YNL113W | RPC19 | DNA-directed RNA polymerase | 1.4870 | 2.8 | 0.0030432 |
| YMR316W | DIA1 | unknown | 1.5038 | 2.8 | 0.0101384 |
| YNL110C | NOP15 | unknown | 1.5062 | 2.8 | 0.0020962 |
| YNL107W | YAF9 | unknown | 1.5532 | 2.9 | 0.0088094 |
| YGR242W | | | 1.5540 | 2.9 | 0.0499112 |
| YMR090W | | unknown | 1.5587 | 2.9 | 0.0062352 |
| YBR004C | GPI18 | mannosyltransferase | 1.5660 | 3.0 | 0.0095197 |
| YBR005W | RCR1 | unknown | 1.5703 | 3.0 | 0.0053983 |
| YJL088W | ARG3 | ornithine carbamoyltransferase | 1.5882 | 3.0 | 0.0012621 |
| YIL003W | CFD1 | ATPase | 1.5893 | 3.0 | 0.0063768 |
| YFL050C | ALR2 | di-, tri-valent inorganic cation transporter | 1.5905 | 3.0 | 0.0061501 |
| YOL142W | RRP40 | 3'-5'-exoribonuclease | 1.6503 | 3.1 | 0.0068724 |
| YOR303W | CPA1 | carbamoyl-phosphate synthase (glutamine-hydrolyzing) | 1.6518 | 3.1 | 0.0501503 |
| YML004C | GLO1 | lactoylglytathione lyase | 1.6558 | 3.2 | 0.0002869 |
| YDR502C | SAM2 | methionine adenosyltransferase | 1.6602 | 3.2 | 0.0007113 |
| YHR137W | ARO9 | aromatic-amino-acid transaminase | 1.6668 | 3.2 | 0.0001437 |
| YDR070C | FMP16 | unknown | 1.6943 | 3.2 | 0.0051151 |
| YMR107W | SPG4 | unknown | 1.6987 | 3.2 | 0.0025338 |
| YER032W | FIR1 | unknown | 1.7165 | 3.3 | 0.003199 |
| YEL071W | DLD3 | D-lactate dehydrogenase (cytochrome) | 1.7180 | 3.3 | 0.0003916 |
| YBR117C | TKL2 | transketolase | 1.7185 | 3.3 | 0.0006054 |
| YDL168W | SFA1 | alcohol dehydrogenase | 1.7803 | 3.4 | 0.0011891 |
| YLR164W | | unknown | 1.8252 | 3.5 | 0.0058369 |
| YGR088W | CTT1 | catalase | 1.8363 | 3.6 | 3.037E-05 |
| YNL193W | | unknown | 1.8413 | 3.6 | 0.003302 |
| YMR176W | ECM5 | unknown | 1.8672 | 3.6 | 0.0032719 |
| YGR243W | FMP43 | unknown | 1.8707 | 3.7 | 0.0098761 |
| YML005W | TRM12 | S-adenosylmethionine-dependent methyltransferase | 1.8733 | 3.7 | 0.0023663 |
| YMR174C | PAI3 | endopeptidase inhibitor | 1.8782 | 3.7 | 0.00829 |
| YNL075W | IMP4 | rRNA primary transcript binding | 1.8830 | 3.7 | 0.0138137 |
| YOR031W | CRS5 | copper ion binding | 1.9465 | 3.9 | 0.0019572 |
| YGL037C | PNC1 | nicotinamidase | 1.9810 | 3.9 | 0.000148 |
| YHR070W | TRM5 | tRNA (guanine) methyltransferase | 1.9837 | 4.0 | 0.0174053 |
| YJL171C | | unknown | 2.0557 | 4.2 | 0.122607 |
| YLR194C | | structural constituent of cell wall | 2.0742 | 4.2 | 0.0058299 |
| YML131W | | unknown | 2.0768 | 4.2 | 0.0002212 |
| YHR034C | PIH1 | unknown | 2.0827 | 4.2 | 0.0055576 |
| YOR100C | CRC1 | carnitine:acyl carnitine antiporter | 2.0887 | 4.3 | 0.0012696 |
| YMR173W-A | | | 2.1310 | 4.4 | 0.0017235 |
| YHR162W | | unknown | 2.2008 | 4.6 | 0.0005867 |
| YDR039C | ENA2 | ATPase , coupled to transmembrane movement of ions, phosph | 2.2312 | 4.7 | 0.0032162 |
| YMR175W | SIP18 | phospholipid binding | 2.2352 | 4.7 | 0.0039071 |
| YHR056C | RSC30 | DNA binding | 2.2843 | 4.9 | 0.0044164 |
| YDR040C | ENA1 | ATPase , coupled to transmembrane movement of ions, phosph | 2.3302 | 5.0 | 0.008578 |
| YDL023C | | | 2.3502 | 5.1 | 0.0038876 |
| YBR116C | | | 2.3518 | 5.1 | 0.0050972 |
| YNL036W | NCE103 | carbonate dehydratase | 2.3613 | 5.1 | 0.0071273 |
| YIL117C | PRM5 | unknown | 2.4263 | 5.4 | 0.0007718 |
| YHR103W | SBE22 | unknown | 2.4640 | 5.5 | 0.0006885 |
| YOR107W | RGS2 | GTPase activator | 2.4728 | 5.6 | 0.0009944 |
| YCR010C | ADY2 | ammonium transporter | 2.4952 | 5.6 | 0.0004029 |
| YFL021W | GAT1 | specific RNA polymerase II transcription factor | 2.5605 | 5.9 | 0.0012964 |
| YNR014W | | unknown | 2.5733 | 6.0 | 0.0038867 |
| YFL029C | CAK1 | cyclin-dependent protein kinase | 2.6262 | 6.2 | 0.0004148 |
| YDL022W | GPD1 | glycerol-3-phosphate dehydrogenase (NAD+) | 2.7333 | 6.6 | 3.523E-05 |
| YPL223C | GRE1 | unknown | 2.7528 | 6.7 | 0.0010662 |
| YJL106W | IME2 | protein kinase | 2.7968 | 6.9 | 0.0002579 |
| YGL039W | | oxidoreductase | 2.8945 | 7.4 | 0.0048832 |
| YGR159C | NSR1 | RNA binding | 2.9822 | 7.9 | 0.0061362 |
| YDR043C | NRG1 | DNA binding | 3.0162 | 8.1 | 2.901E-05 |
| YFL013W-A | | | 3.0492 | 8.3 | 0.0023316 |
| YDL214C | PRR2 | receptor signaling protein serine/threonine kinase | 3.0513 | 8.3 | 0.0042473 |
| YFL014W | HSP12 | unknown | 3.0805 | 8.5 | 0.0051355 |
| YHR139C | SPS100 | unknown | 3.1183 | 8.7 | 0.0352836 |
| YHR033W | | unknown | 3.2067 | 9.2 | 0.0006293 |
| YGL157W | | oxidoreductase | 3.4043 | 10.6 | 0.0032501 |
| YOL150C | | | 3.4392 | 10.8 | 0.0075217 |
| YBR281C | | unknown | 3.5735 | 11.9 | 0.0061927 |
| YER062C | HOR2 | glycerol-1-phosphatase | 3.6217 | 12.3 | 0.0010461 |

Table 3: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|-------------------------------------|----------------------------|-------------|-----------|
| YHR087W | | unknown | 3.6385 | 12.5 | 0.0004163 |
| YIL053W | RHR2 | glycerol-1-phosphatase | 3.8072 | 14.0 | 0.0022372 |
| YMR169C | ALD3 | 3-chloroalyl aldehyde dehydrogenase | 3.8642 | 14.6 | 0.0011308 |
| YHR055C | CUP1-2 | copper ion binding | 3.9318 | 15.3 | 0.0027359 |
| YHR053C | CUP1-1 | copper ion binding | 3.9465 | 15.4 | 0.0026594 |
| YDR536W | STL1 | transporter | 4.3728 | 20.7 | 0.0003767 |
| YOL151W | GRE2 | oxidoreductase | 4.6408 | 24.9 | 0.0013843 |

Table 4: Raw microarray data for fermentation day five.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YDR444W | | unknown | -5.3392 | -40.5 | 0.051763 |
| YAL048C | GEM1 | GTPase | -4.5062 | -22.7 | 0.0344011 |
| YCR045C | | serine-type peptidase | -4.2547 | -19.1 | 0.0440621 |
| YLR382C | NAM2 | mRNA binding | -4.2263 | -18.7 | 0.0289843 |
| YGR204W | ADE3 | formate-tetrahydrofolate ligase | -4.1873 | -18.2 | 0.0548038 |
| YMR212C | EFR3 | unknown | -3.9447 | -15.4 | 0.0062186 |
| YBR229C | ROT2 | alpha-glucosidase | -3.9010 | -14.9 | 0.026299 |
| YGR098C | ESP1 | cysteine-type endopeptidase | -3.7858 | -13.8 | 0.0679902 |
| YGR254W | ENO1 | phosphopyruvate hydratase | -3.7653 | -13.6 | 0.0074388 |
| YKR002W | PAP1 | polynucleotide adenyltransferase | -3.7153 | -13.1 | 0.0256818 |
| YBL101C | ECM21 | unknown | -3.6018 | -12.1 | 0.0211561 |
| YLR410W | VIP1 | unknown | -3.5778 | -11.9 | 0.0305307 |
| YGR087C | PDC6 | pyruvate decarboxylase | -3.3335 | -10.1 | 0.0353946 |
| YGL253W | HXK2 | hexokinase | -3.2457 | -9.5 | 0.0435592 |
| YBL066C | SEF1 | unknown | -3.1123 | -8.6 | 0.207726 |
| YGL243W | TAD1 | tRNA specific adenosine deaminase | -3.0443 | -8.2 | 0.0676653 |
| YHR092C | HXT4 | glucose transporter | -2.9675 | -7.8 | 0.0073077 |
| Q0255 | | unknown | -2.9470 | -7.7 | 0.0239738 |
| YCR041W | | | -2.9003 | -7.5 | 0.0201269 |
| YLR304C | ACO1 | aconitate hydratase | -2.8653 | -7.3 | 0.0766614 |
| YDR385W | EFT2 | translation elongation factor | -2.7742 | -6.8 | 0.076096 |
| YGR192C | TDH3 | glyceraldehyde-3-phosphate dehydrogenase | -2.7288 | -6.6 | 0.0159984 |
| YFR053C | HXK1 | hexokinase | -2.7267 | -6.6 | 0.0465588 |
| YMR302C | YME2 | exonuclease | -2.7135 | -6.6 | 0.0355041 |
| YBR115C | LYS2 | L-aminoadipate-semialdehyde dehydrogenase | -2.6073 | -6.1 | 0.0422329 |
| YOR306C | MCH5 | transporter | -2.5742 | -6.0 | 0.0580967 |
| YCR025C | | | -2.5660 | -5.9 | 0.0501329 |
| YKL171W | | protein kinase | -2.5295 | -5.8 | 0.023167 |
| YCR044C | PER1 | unknown | -2.5255 | -5.8 | 0.0482678 |
| YDR249C | | unknown | -2.4080 | -5.3 | 0.0256315 |
| YOR375C | GDH1 | glutamate dehydrogenase (NADP+) | -2.3998 | -5.3 | 0.0137396 |
| YBR084W | MIS1 | formate-tetrahydrofolate ligase | -2.3965 | -5.3 | 0.0266945 |
| YMR303C | ADH2 | alcohol dehydrogenase | -2.2927 | -4.9 | 0.0428944 |
| YHR094C | HXT1 | glucose transporter | -2.2070 | -4.6 | 0.20141 |
| YPR035W | GLN1 | glutamate-ammonia ligase | -2.0923 | -4.3 | 0.0213266 |
| YHR096C | HXT5 | glucose transporter | -2.0707 | -4.2 | 0.0139458 |
| YIL026C | IRR1 | protein binding | -2.0672 | -4.2 | 0.0960565 |
| YBR127C | VMA2 | hydrogen-transporting ATPase | -2.0652 | -4.2 | 0.0232156 |
| YDR516C | EMI2 | unknown | -2.0278 | -4.1 | 0.0430584 |
| YHR174W | ENO2 | phosphopyruvate hydratase | -2.0000 | -4.0 | 0.0394261 |
| YLR044C | PDC1 | pyruvate decarboxylase | -1.9802 | -3.9 | 0.0226271 |
| YBL075C | SSA3 | ATPase | -1.9228 | -3.8 | 0.022582 |
| YBR166C | TYR1 | prephenate dehydrogenase | -1.9082 | -3.8 | 0.0565102 |
| YLR454W | FMP27 | unknown | -1.8977 | -3.7 | 0.018047 |
| YHR181W | SVP26 | protein binding | -1.8647 | -3.6 | 0.082482 |
| YNL072W | RNH201 | ribonuclease H | -1.8630 | -3.6 | 0.0426152 |
| YMR083W | ADH3 | alcohol dehydrogenase | -1.8503 | -3.6 | 0.0911022 |
| YDR247W | VHS1 | protein kinase | -1.8400 | -3.6 | 0.0152159 |
| YMR163C | | unknown | -1.8355 | -3.6 | 0.0534559 |
| YEL060C | PRB1 | serine-type endopeptidase | -1.8322 | -3.6 | 0.0182008 |
| YMR297W | PRC1 | carboxypeptidase C | -1.8112 | -3.5 | 0.0035305 |
| YHL048W | COS8 | unknown | -1.8012 | -3.5 | 0.0344361 |
| YDR246W | TRS23 | unknown | -1.7747 | -3.4 | 0.0161081 |
| YJL034W | KAR2 | ATPase | -1.7743 | -3.4 | 0.0532092 |
| YFL062W | COS4 | unknown | -1.7330 | -3.3 | 0.047555 |
| YNL209W | SSB2 | ATPase | -1.6767 | -3.2 | 0.0339435 |
| YOL086C | ADH1 | alcohol dehydrogenase | -1.6753 | -3.2 | 0.0227968 |
| YML132W | COS3 | protein binding | -1.6287 | -3.1 | 0.0225979 |
| YCL024W | KCC4 | protein kinase | -1.6078 | -3.0 | 0.11384 |
| YJR016C | ILV3 | dihydroxy-acid dehydratase | -1.6025 | -3.0 | 0.0231506 |
| YMR125W | STO1 | mRNA binding | -1.5743 | -3.0 | 0.0507185 |
| YBL099W | ATP1 | hydrogen-transporting ATP synthase | -1.5735 | -3.0 | 0.0448278 |
| YKL103C | LAP4 | aminopeptidase I | -1.5043 | -2.8 | 0.0028877 |
| YJR121W | ATP2 | hydrogen-transporting ATP synthase | -1.5003 | -2.8 | 0.0009516 |
| YGR295C | COS6 | unknown | -1.4972 | -2.8 | 0.0492923 |
| YDR129C | SAC6 | protein binding | -1.4763 | -2.8 | 0.0492033 |
| YOR007C | SGT2 | unknown | -1.4702 | -2.8 | 0.0545219 |
| YGR282C | BGL2 | glucan 1,3-beta-glucosidase | -1.4538 | -2.7 | 0.0393614 |
| YAL037C-B | | | -1.4488 | -2.7 | 0.0424666 |

Table 4: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|---------|-----------|---|----------------------------|-------------|-----------|
| YNL336W | COS1 | unknown | -1.4452 | -2.7 | 0.0542555 |
| YGR248W | SOL4 | 6-phosphogluconolactonase | -1.4103 | -2.7 | 0.0447699 |
| YNL007C | SIS1 | unfolded protein binding | -1.4067 | -2.7 | 0.035954 |
| YNL055C | POR1 | voltage-gated ion-selective channel | -1.4002 | -2.6 | 0.0474275 |
| YBR128C | ATG14 | unknown | -1.3803 | -2.6 | 0.0351967 |
| YJL052W | TDH1 | glyceraldehyde-3-phosphate dehydrogenase | -1.3770 | -2.6 | 0.010066 |
| YDR432W | NPL3 | mRNA binding | -1.3710 | -2.6 | 0.0297018 |
| YFL039C | ACT1 | structural constituent of cytoskeleton | -1.3562 | -2.6 | 0.0388893 |
| YCR029C | | | -1.3558 | -2.6 | 0.014576 |
| YJR009C | TDH2 | glyceraldehyde-3-phosphate dehydrogenase | -1.3525 | -2.6 | 0.0546887 |
| YAL062W | GDH3 | glutamate dehydrogenase | -1.3248 | -2.5 | 0.103498 |
| YHR126C | | unknown | -1.3195 | -2.5 | 0.0325986 |
| YJL160C | | unknown | -1.2820 | -2.4 | 0.0105725 |
| YPL240C | HSP82 | ATPase | -1.2735 | -2.4 | 0.0281381 |
| YML028W | TSA1 | thioredoxin peroxidase | -1.2655 | -2.4 | 0.0325153 |
| YHR214W | | unknown | -1.2520 | -2.4 | 0.0587032 |
| YBL045C | COR1 | ubiquinol-cytochrome-c reductase | -1.2448 | -2.4 | 0.035604 |
| YCR004C | YCP4 | electron carrier | -1.2435 | -2.4 | 0.0344376 |
| YDR032C | PST2 | unknown | -1.2425 | -2.4 | 0.037142 |
| YKL117W | SBA1 | chaperone binding | -1.2380 | -2.4 | 0.0399051 |
| YLR257W | | unknown | -1.2313 | -2.3 | 0.0924239 |
| YDL029W | ARP2 | structural constituent of cytoskeleton* | -1.2243 | -2.3 | 0.178642 |
| YMR186W | HSC82 | unfolded protein binding | -1.2205 | -2.3 | 0.0007939 |
| YOR027W | STI1 | unfolded protein binding | -1.1647 | -2.2 | 0.0468117 |
| YBR144C | | | -1.1547 | -2.2 | 0.0182665 |
| YPR033C | HTS1 | histidine-tRNA ligase | -1.1540 | -2.2 | 0.0207082 |
| YCL042W | | unknown | -1.1455 | -2.2 | 0.0527729 |
| YML100W | TSL1 | enzyme regulator | -1.1102 | -2.2 | 0.0200636 |
| YLR258W | GSY2 | glycogen (starch) synthase | -1.1092 | -2.2 | 0.0075418 |
| YPR080W | TEF1 | translation elongation factor | -1.0617 | -2.1 | 0.0257002 |
| YPR113W | PIS1 | CDP-diaclyglycerol-inositol 3-phosphatidyltransferase | -1.0612 | -2.1 | 0.0497974 |
| YJL079C | PRY1 | unknown | -1.0538 | -2.1 | 0.050628 |
| YGL076C | RPL7A | structural constituent of ribosome | -1.0323 | -2.0 | 0.0160039 |
| YBL001C | ECM15 | unknown | -1.0195 | -2.0 | 0.0485627 |
| YLR149C | | unknown | -1.0177 | -2.0 | 0.0998473 |
| YDR453C | TSA2 | thioredoxin peroxidase | -1.0160 | -2.0 | 0.0330834 |
| YHR051W | COX6 | cytochrome-c oxidase | -1.0148 | -2.0 | 0.0169114 |
| YDR322W | MRPL35 | structural constituent of ribosome | -0.9937 | -2.0 | 0.0511027 |
| YAL002W | VPS8 | unknown | -0.9903 | -2.0 | 0.401723 |
| YNL037C | IDH1 | isocitrate dehydrogenase (NAD ⁺) | -0.9893 | -2.0 | 0.0766296 |
| YKL163W | PIR3 | structural constituent of cell wall | -0.9827 | -2.0 | 0.0361575 |
| YPL037C | EGD1 | unfolded protein binding | -0.9800 | -2.0 | 0.0541988 |
| YLR109W | AHP1 | thioredoxin peroxidase | -0.9773 | -2.0 | 0.04681 |
| YER053C | PIC2 | inorganic phosphate transporter | -0.9753 | -2.0 | 0.012701 |
| YBR218C | PYC2 | pyruvate carboxylase | -0.9668 | -2.0 | 0.067544 |
| YBR286W | APE3 | aminopeptidase | -0.9597 | -1.9 | 0.094082 |
| YKL067W | YNK1 | nucleoside diphosphate kinase | -0.9515 | -1.9 | 0.0193619 |
| YDR163W | CWC15 | unknown | -0.9508 | -1.9 | 0.0245781 |
| YIL018W | RPL2B | structural constituent of ribosome | -0.9340 | -1.9 | 0.0347385 |
| YGR180C | RNR4 | ribonucleoside-diphosphate reductase | -0.9320 | -1.9 | 0.027898 |
| YDR157W | | | -0.9258 | -1.9 | 0.0415326 |
| YNL208W | | unknown | -0.9232 | -1.9 | 0.121175 |
| YAL005C | SSA1 | ATPase | -0.9035 | -1.9 | 0.00993 |
| YBL030C | PET9 | ATP:ADP antiporter | -0.9015 | -1.9 | 0.24093 |
| YIL094C | LYS12 | homoisocitrate dehydrogenase | -0.8955 | -1.9 | 0.0934259 |
| YJL129C | TRK1 | potassium ion transporter | -0.8778 | -1.8 | 0.0894357 |
| YCL009C | ILV6 | enzyme regulator | -0.8747 | -1.8 | 0.0663351 |
| YCL064C | CHA1 | L-serine ammonia-lyase | -0.8688 | -1.8 | 0.206763 |
| YHR219W | | unknown | -0.8632 | -1.8 | 0.0186118 |
| YNL064C | YDJ1 | chaperone regulator | -0.8600 | -1.8 | 0.21054 |
| YKL013C | ARC19 | structural molecule | -0.8495 | -1.8 | 0.178514 |
| YMR233W | | unknown | -0.8493 | -1.8 | 0.0104045 |
| YIL138C | TPM2 | actin lateral binding | -0.8487 | -1.8 | 0.0239481 |
| YJL223C | PAU1 | unknown | -0.8458 | -1.8 | 0.01429 |
| YBR079C | RPG1 | translation initiation factor | -0.8167 | -1.8 | 0.0203147 |
| YJR103W | URA8 | CTP synthase | -0.7887 | -1.7 | 0.0035963 |
| YEL027W | CUP5 | hydrogen ion transporter | -0.7868 | -1.7 | 0.0142962 |
| YMR260C | TIF11 | translation initiation factor | -0.7867 | -1.7 | 0.0541752 |
| YBR203W | COS111 | unknown | -0.7847 | -1.7 | 0.0256541 |

Table 4: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YJL159W | HSP150 | structural constituent of cell wall | -0.7770 | -1.7 | 0.0324755 |
| YEL077C | | helicase | -0.7742 | -1.7 | 0.0517198 |
| YAR020C | PAU7 | unknown | -0.7555 | -1.7 | 0.0127277 |
| YLR441C | RPS1A | structural constituent of ribosome | -0.7533 | -1.7 | 0.0517316 |
| YLR388W | RPS29A | structural constituent of ribosome | -0.7532 | -1.7 | 0.502622 |
| YHL033C | RPL8A | structural constituent of ribosome | -0.7343 | -1.7 | 0.0251955 |
| YPR108W | RPN7 | structural molecule | -0.7202 | -1.6 | 0.186524 |
| YOR185C | GSP2 | GTPase | -0.7197 | -1.6 | 0.0402039 |
| YIR043C | | unknown | -0.7132 | -1.6 | 0.0508439 |
| YKL152C | GPM1 | phosphoglycerate mutase | -0.7105 | -1.6 | 0.0322486 |
| YKL058W | TOA2 | general RNA polymerase II transcription factor | -0.6935 | -1.6 | 0.0162936 |
| YHL050C | | helicase | -0.6908 | -1.6 | 0.042141 |
| YDL137W | ARF2 | GTPase | -0.6883 | -1.6 | 0.0986587 |
| YOR332W | VMA4 | hydrogen-transporting ATPase | -0.6848 | -1.6 | 0.0515079 |
| YDR086C | SSS1 | protein transporter | -0.6682 | -1.6 | 0.152408 |
| YKL056C | TMA19 | unknown | -0.6680 | -1.6 | 0.130941 |
| YLR264W | RPS28B | structural constituent of ribosome | -0.6642 | -1.6 | 0.0213802 |
| YOR167C | RPS28A | structural constituent of ribosome | -0.6625 | -1.6 | 0.0292391 |
| YAL056W | GPB2 | signal transducer | -0.6580 | -1.6 | 0.0421383 |
| YAL013W | DEP1 | transcription regulator | -0.6548 | -1.6 | 0.0955915 |
| YBR237W | PRP5 | RNA splicing factor | -0.6533 | -1.6 | 0.0745361 |
| YGR118W | RPS23A | structural constituent of ribosome | -0.6513 | -1.6 | 0.0279966 |
| YKL160W | ELF1 | RNA polymerase II transcription elongation factor | -0.6422 | -1.6 | 0.0492036 |
| YIR037W | HYR1 | thioredoxin peroxidase | -0.6337 | -1.6 | 0.0036256 |
| YIL011W | TIR3 | unknown | -0.6213 | -1.5 | 0.0372393 |
| YCL018W | LEU2 | 3-isopropylmalate dehydrogenase | -0.6213 | -1.5 | 0.01219 |
| YNL079C | TPM1 | actin lateral binding | -0.6210 | -1.5 | 0.0307504 |
| YBR170C | NPL4 | unknown | -0.6182 | -1.5 | 0.0744633 |
| YBL002W | HTB2 | DNA binding | -0.6155 | -1.5 | 0.294039 |
| YKL006W | RPL14A | structural constituent of ribosome | -0.6150 | -1.5 | 0.0006214 |
| YCR032W | BPH1 | unknown | -0.6087 | -1.5 | 0.259111 |
| YKR042W | UTH1 | unknown | -0.6080 | -1.5 | 0.173422 |
| YBR301W | DAN3 | unknown | -0.6070 | -1.5 | 0.0276601 |
| YBR263W | SHM1 | glycine hydroxymethyltransferase | -0.6060 | -1.5 | 0.0027566 |
| YBR121C | GRS1 | glycine-tRNA ligase | -0.6042 | -1.5 | 0.0469617 |
| YHR008C | SOD2 | manganese superoxide dismutase | -0.5990 | -1.5 | 0.0392052 |
| YOR215C | | unknown | -0.5953 | -1.5 | 0.0125881 |
| YLR238W | FAR10 | unknown | -0.5887 | -1.5 | 0.0507794 |
| YLR325C | RPL38 | structural constituent of ribosome | -0.5853 | -1.5 | 0.0236712 |
| YIL155C | GUT2 | glycerol-3-phosphate dehydrogenase | -0.5823 | -1.5 | 0.210418 |
| YIR041W | PAU15 | unknown | -0.5755 | -1.5 | 0.0400524 |
| YBR164C | ARL1 | GTPase | -0.5698 | -1.5 | 0.0646627 |
| YEL054C | RPL12A | structural constituent of ribosome | -0.5673 | -1.5 | 0.139228 |
| YBR010W | HHT1 | DNA binding | -0.5668 | -1.5 | 0.053004 |
| YJL177W | RPL17B | structural constituent of ribosome | -0.5557 | -1.5 | 0.0615107 |
| YBR082C | UBC4 | ubiquitin conjugating enzyme | -0.5533 | -1.5 | 0.277941 |
| YDR033W | MRH1 | unknown | -0.5392 | -1.5 | 0.043695 |
| YLR227W-A | | | -0.5375 | -1.5 | 0.0497666 |
| YIL142W | CCT2 | unfolded protein binding | -0.5350 | -1.4 | 0.0500548 |
| YBR221C | PDB1 | pyruvate dehydrogenase (acetyl-transferring) | -0.5342 | -1.4 | 0.0522371 |
| YCR047C | BUD23 | S-adenosylmethionine-dependent methyltransferase | -0.5338 | -1.4 | 0.157489 |
| YER057C | HMF1 | unknown | -0.5293 | -1.4 | 0.0478953 |
| YKR094C | RPL40B | structural constituent of ribosome | -0.5022 | -1.4 | 0.148783 |
| YJL158C | CIS3 | structural constituent of cell wall | -0.5017 | -1.4 | 0.0727102 |
| YBR081C | SPT7 | structural molecule | -0.4840 | -1.4 | 0.365906 |
| YBR067C | TIP1 | structural constituent of cell wall* | -0.4827 | -1.4 | 0.251577 |
| YKL060C | FBA1 | fructose-bisphosphate aldolase | -0.4808 | -1.4 | 0.278732 |
| YBR126C | TPS1 | alpha, alpha-trehalose-phosphate synthase (UDP-forming) | -0.4728 | -1.4 | 0.0825319 |
| YPL090C | RPS6A | structural constituent of ribosome | -0.4703 | -1.4 | 0.0035684 |
| YKL085W | MDH1 | L-malate dehydrogenase | -0.4592 | -1.4 | 0.0489727 |
| YHR135C | YCK1 | casein kinase I | -0.4468 | -1.4 | 0.413325 |
| YER102W | RPS8B | structural constituent of ribosome | -0.4345 | -1.4 | 0.0049872 |
| YLR043C | TRX1 | thiol-disulfide exchange intermediate | -0.4305 | -1.3 | 0.0060027 |
| YML045W | | | -0.4202 | -1.3 | 0.0193989 |
| YEL049W | PAU2 | unknown | -0.4198 | -1.3 | 0.0016843 |
| YCR104W | PAU3 | unknown | -0.4102 | -1.3 | 0.0137731 |
| YLR216C | CPR6 | unfolded protein binding | -0.4013 | -1.3 | 0.0084931 |
| YGR037C | ACB1 | long-chain fatty acid transporter | -0.3985 | -1.3 | 0.0046039 |
| YJR073C | OPI3 | phosphatidyl-N-methylethanolamine N-methyltransferase | -0.3958 | -1.3 | 0.0340325 |

Table 4: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YIL176C | PAU14 | unknown | -0.3848 | -1.3 | 0.16288 |
| YDR201W | SPC19 | structural constituent of cytoskeleton | -0.3767 | -1.3 | 0.475063 |
| YBR234C | ARC40 | structural constituent of cytoskeleton | -0.3733 | -1.3 | 0.0467256 |
| YBR255W | | unknown | -0.3728 | -1.3 | 0.0621633 |
| YMR190C | SGS1 | ATP-dependent DNA helicase | -0.3702 | -1.3 | 0.215257 |
| YDR500C | RPL37B | structural constituent of ribosome | -0.3658 | -1.3 | 0.0677628 |
| YBR208C | DUR1,2 | allophanate hydrolase | -0.3618 | -1.3 | 0.14266 |
| YBR094W | | unknown | -0.3533 | -1.3 | 0.0108237 |
| YDR119W | | unknown | -0.3513 | -1.3 | 0.0295702 |
| YHR192W | | unknown | -0.3432 | -1.3 | 0.195677 |
| YBR080C | SEC18 | ATPase | -0.3407 | -1.3 | 0.226605 |
| YBL027W | RPL19B | structural constituent of ribosome | -0.3385 | -1.3 | 0.525091 |
| YBR224W | | | -0.3185 | -1.2 | 0.0180677 |
| YBL113C | | helicase | -0.2948 | -1.2 | 0.191124 |
| YBR191W | RPL21A | structural constituent of ribosome | -0.2843 | -1.2 | 0.0247335 |
| YIL051C | MMF1 | unknown | -0.2795 | -1.2 | 0.473004 |
| YDR382W | RPP2B | structural constituent of ribosome | -0.2795 | -1.2 | 0.0472885 |
| YNL160W | YGP1 | unknown | -0.2737 | -1.2 | 0.200024 |
| YAL068C | PAU8 | unknown | -0.2683 | -1.2 | 0.0073871 |
| YHR183W | GND1 | phosphogluconate dehydrogenase (decarboxylating) | -0.2590 | -1.2 | 0.742296 |
| YNL161W | CBK1 | protein kinase | -0.2577 | -1.2 | 0.656745 |
| YLR340W | RPP0 | structural constituent of ribosome | -0.2555 | -1.2 | 0.220982 |
| YCR053W | THR4 | threonine synthase | -0.2548 | -1.2 | 0.0314283 |
| YLL066W-A | | | -0.2465 | -1.2 | 0.68185 |
| YBR012W-A | | | -0.2445 | -1.2 | 0.209382 |
| YBR031W | RPL4A | structural constituent of ribosome | -0.2412 | -1.2 | 0.787372 |
| YGR148C | RPL24B | structural constituent of ribosome | -0.2347 | -1.2 | 0.257877 |
| YJR028W | | | -0.2337 | -1.2 | 0.0340705 |
| YOR096W | RPS7A | structural constituent of ribosome | -0.2308 | -1.2 | 0.0456217 |
| YBR125C | PTC4 | protein phosphatase type 2C | -0.2285 | -1.2 | 0.179713 |
| YMR264W | CUE1 | protein binding | -0.2262 | -1.2 | 0.0467298 |
| YER068W | MOT2 | 3'-5'-exoribonuclease | -0.2248 | -1.2 | 0.0062249 |
| YGL095C | VPS45 | unfolded protein binding | -0.2145 | -1.2 | 0.0225472 |
| YBR212W | NGR1 | RNA binding | -0.2125 | -1.2 | 0.71913 |
| YBR072W | HSP26 | unfolded protein binding | -0.2118 | -1.2 | 0.290927 |
| YDL082W | RPL13A | structural constituent of ribosome | -0.2072 | -1.2 | 0.291089 |
| YBR086C | IST2 | unknown | -0.2028 | -1.2 | 0.374258 |
| YLR333C | RPS25B | structural constituent of ribosome | -0.2002 | -1.1 | 0.0631351 |
| YDR202C | RAV2 | unknown | -0.1908 | -1.1 | 0.695319 |
| YAL035W | FUN12 | GTPase | -0.1877 | -1.1 | 0.261907 |
| YJL020C | BBC1 | myosin I binding | -0.1867 | -1.1 | 0.0357772 |
| YBR069C | TAT1 | amino acid transporter | -0.1833 | -1.1 | 0.825265 |
| YJL210W | PEX2 | protein binding | -0.1742 | -1.1 | 0.0424187 |
| YBR095C | RXT2 | unknown | -0.1717 | -1.1 | 0.432364 |
| YBR009C | HHF1 | DNA binding | -0.1688 | -1.1 | 0.0446996 |
| YLR185W | RPL37A | structural constituent of ribosome | -0.1630 | -1.1 | 0.0507136 |
| YCR028C | FEN2 | pantothenate transporter | -0.1535 | -1.1 | 0.835791 |
| YBR148W | YSW1 | unknown | -0.1465 | -1.1 | 0.445584 |
| YJL111W | CCT7 | unfolded protein binding | -0.1455 | -1.1 | 0.485339 |
| YJL174W | KRE9 | unknown | -0.1437 | -1.1 | 0.515636 |
| YDR321W | ASP1 | asparaginase | -0.1402 | -1.1 | 0.0219457 |
| YDR329C | PEX3 | protein binding | -0.1383 | -1.1 | 0.0065111 |
| YBR111C | YSA1 | phosphoribosyl-ATP diphosphatase | -0.1202 | -1.1 | 0.46628 |
| YDR080W | VPS41 | Rab guanyl-nucleotide exchange factor | -0.1170 | -1.1 | 0.470536 |
| YCL025C | AGP1 | amino acid transporter | -0.1077 | -1.1 | 0.451172 |
| YBR280C | | unknown | -0.1060 | -1.1 | 0.458173 |
| YBR053C | | unknown | -0.0850 | -1.1 | 0.567644 |
| YLR294C | | | -0.0837 | -1.1 | 0.554822 |
| YGL135W | RPL1B | structural constituent of ribosome | -0.0812 | -1.1 | 0.605322 |
| YDR542W | PAU10 | unknown | -0.0712 | -1.1 | 0.20482 |
| YBR302C | COS2 | unknown | -0.0512 | -1.0 | 0.788207 |
| YBR008C | FLR1 | multidrug transporter | -0.0457 | -1.0 | 0.776315 |
| YJL070C | | unknown | -0.0335 | -1.0 | 0.862868 |
| YBR205W | KTR3 | mannosyltransferase | 0.0028 | 1.0 | 0.987206 |
| YAL021C | CCR4 | 3'-5'-exoribonuclease | 0.0102 | 1.0 | 0.954663 |
| YMR143W | RPS16A | structural constituent of ribosome | 0.0120 | 1.0 | 0.965284 |
| YCRX13W | | | 0.0140 | 1.0 | 0.949018 |
| YER177W | BMH1 | protein binding | 0.0255 | 1.0 | 0.626577 |
| YMR242C | RPL20A | structural constituent of ribosome | 0.0318 | 1.0 | 0.565757 |

Table 4: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|--|----------------------------|-------------|-----------|
| YOR182C | RPS30B | structural constituent of ribosome | 0.0530 | 1.0 | 0.925237 |
| YGL189C | RPS26A | structural constituent of ribosome | 0.0543 | 1.0 | 0.713676 |
| YER067W | | unknown | 0.0617 | 1.0 | 0.450352 |
| YBR259W | | unknown | 0.0647 | 1.0 | 0.738226 |
| YDR025W | RPS11A | structural constituent of ribosome | 0.0705 | 1.1 | 0.41071 |
| YER011W | TIR1 | structural constituent of cell wall | 0.0707 | 1.1 | 0.67071 |
| YDL136W | RPL35B | structural constituent of ribosome | 0.0735 | 1.1 | 0.607779 |
| YBR262C | FMP51 | unknown | 0.0795 | 1.1 | 0.599434 |
| YCL020W | | | 0.1157 | 1.1 | 0.587526 |
| YIL043C | CBR1 | cytochrome-b5 reductase | 0.1212 | 1.1 | 0.568691 |
| YDR377W | ATP17 | hydrogen-transporting ATP synthase | 0.1287 | 1.1 | 0.0463994 |
| YIL074C | SER33 | phosphoglycerate dehydrogenase | 0.1287 | 1.1 | 0.0528452 |
| YHR190W | ERG9 | farnesyl-diphosphate farnesyltransferase | 0.1307 | 1.1 | 0.758866 |
| YOR285W | | unknown | 0.1383 | 1.1 | 0.548149 |
| YDR273W | DON1 | unknown | 0.1590 | 1.1 | 0.297698 |
| YDR473C | PRP3 | RNA splicing factor | 0.1812 | 1.1 | 0.0011975 |
| YLR395C | COX8 | cytochrome-c oxidase | 0.1837 | 1.1 | 0.0267468 |
| YBL092W | RPL32 | structural constituent of ribosome | 0.1968 | 1.1 | 0.283901 |
| YNL149C | PGA2 | unknown | 0.1978 | 1.1 | 0.447352 |
| YJL225C | | helicase | 0.2048 | 1.2 | 0.0025009 |
| YDL075W | RPL31A | structural constituent of ribosome | 0.2070 | 1.2 | 0.477447 |
| YJL021C | | | 0.2092 | 1.2 | 0.0208446 |
| YMR022W | QRI8 | ubiquitin conjugating enzyme | 0.2112 | 1.2 | 0.0129973 |
| YLR193C | | unknown | 0.2177 | 1.2 | 0.0363107 |
| YPR020W | ATP20 | structural molecule | 0.2223 | 1.2 | 0.0425931 |
| YER019C-A | SBH2 | protein transporter | 0.2225 | 1.2 | 0.566916 |
| YIL105C | SLM1 | phosphoinositide binding | 0.2287 | 1.2 | 0.015514 |
| YDR154C | | | 0.2290 | 1.2 | 0.020685 |
| YHR104W | GRE3 | aldo-keto reductase | 0.2337 | 1.2 | 0.0392109 |
| YNL099C | OCA1 | protein tyrosine phosphatase | 0.2358 | 1.2 | 0.502156 |
| YCL002C | | unknown | 0.2395 | 1.2 | 0.031586 |
| YBR135W | CKS1 | protein kinase activator | 0.2397 | 1.2 | 0.0667429 |
| YOR113W | AZF1 | DNA binding | 0.2483 | 1.2 | 0.051918 |
| YJR048W | CYC1 | electron carrier | 0.2588 | 1.2 | 0.0315101 |
| YOR394W | PAU21 | unknown | 0.2638 | 1.2 | 0.758736 |
| YGL227W | VID30 | unknown | 0.2778 | 1.2 | 0.0204272 |
| YOR310C | NOP58 | unknown | 0.2788 | 1.2 | 0.0046976 |
| YLR158C | ASP3-3 | asparaginase | 0.2813 | 1.2 | 0.261902 |
| YDR145W | TAF12 | general RNA polymerase II transcription factor | 0.2833 | 1.2 | 0.0472565 |
| YAL016W | TPD3 | protein phosphatase type 2A | 0.2882 | 1.2 | 0.216513 |
| YOR359W | VTS1 | RNA binding | 0.2905 | 1.2 | 0.0517269 |
| YJR104C | SOD1 | copper, zinc superoxide dismutase | 0.2927 | 1.2 | 0.261885 |
| YBR238C | | unknown | 0.2982 | 1.2 | 0.0092437 |
| YER056C-A | RPL34A | structural constituent of ribosome | 0.2995 | 1.2 | 0.0129893 |
| YHR146W | CRP1 | DNA binding | 0.3000 | 1.2 | 0.035565 |
| YDR002W | YRB1 | Ran GTPase binding | 0.3002 | 1.2 | 0.433006 |
| YOR042W | CUE5 | unknown | 0.3065 | 1.2 | 0.0020051 |
| YBR172C | SMY2 | unknown | 0.3128 | 1.2 | 0.0021384 |
| YJL136C | RPS21B | structural constituent of ribosome | 0.3163 | 1.2 | 0.0276983 |
| YCR024C-A | PMP1 | enzyme regulator | 0.3173 | 1.2 | 0.152376 |
| YCRX16C | | | 0.3207 | 1.2 | 0.0640786 |
| YBR102C | EXO84 | protein binding | 0.3283 | 1.3 | 0.329939 |
| YCR084C | TUP1 | general transcriptional repressor | 0.3310 | 1.3 | 0.0360097 |
| YBR261C | | S-adenosylmethionine-dependent methyltransferase | 0.3343 | 1.3 | 0.0067372 |
| YHR152W | SPO12 | unknown | 0.3378 | 1.3 | 0.107735 |
| YER080W | FMP29 | unknown | 0.3388 | 1.3 | 0.0069813 |
| YDL124W | | alpha-keto amide reductase | 0.3392 | 1.3 | 0.26327 |
| YNL284C-A | | | 0.3402 | 1.3 | 0.0209582 |
| YDR034C | LYS14 | transcriptional activator | 0.3420 | 1.3 | 0.0443624 |
| YDL134C | PPH21 | protein phosphatase type 2A | 0.3435 | 1.3 | 0.27363 |
| YER109C | FLO8 | specific RNA polymerase II transcription factor | 0.3557 | 1.3 | 0.0110862 |
| YIL068C | SEC6 | protein binding | 0.3567 | 1.3 | 0.452916 |
| YBR268W | MRPL37 | structural constituent of ribosome | 0.3572 | 1.3 | 0.0070668 |
| YJL141C | YAK1 | protein kinase | 0.3653 | 1.3 | 0.112743 |
| YDR134C | | unknown | 0.3703 | 1.3 | 0.177762 |
| YDR510W | SMT3 | protein tag | 0.3722 | 1.3 | 0.0119308 |
| YDR309C | GIC2 | small GTPase regulator | 0.3752 | 1.3 | 0.350099 |
| YAL030W | SNC1 | v-SNARE | 0.3765 | 1.3 | 0.0967859 |
| YLR204W | QRI5 | unknown | 0.3785 | 1.3 | 0.0028322 |

Table 4: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YHR200W | RPN10 | endopeptidase | 0.3908 | 1.3 | 0.0867604 |
| YIL045W | PIG2 | protein phosphatase type 1 regulator | 0.3923 | 1.3 | 0.0317944 |
| YIL052C | RPL34B | structural constituent of ribosome | 0.3963 | 1.3 | 0.0234255 |
| YNL027W | CRZ1 | transcription factor | 0.3967 | 1.3 | 0.010022 |
| YLR093C | NYV1 | v-SNARE | 0.3972 | 1.3 | 0.0210133 |
| YCR042C | TAF2 | general RNA polymerase II transcription factor | 0.3993 | 1.3 | 0.0430635 |
| YBR117C | TKL2 | transketolase | 0.4085 | 1.3 | 0.0265663 |
| YBR071W | | unknown | 0.4103 | 1.3 | 0.0156187 |
| YOR327C | SNC2 | v-SNARE | 0.4108 | 1.3 | 0.0424408 |
| YOR065W | CYT1 | electron transporter | 0.4218 | 1.3 | 0.017113 |
| YNR052C | POP2 | 3'-5'-exoribonuclease | 0.4298 | 1.3 | 0.0398091 |
| YBR090C | | unknown | 0.4330 | 1.4 | 0.117835 |
| YDR261C-C | | | 0.4348 | 1.4 | 0.0394559 |
| YDL147W | RPN5 | unknown | 0.4350 | 1.4 | 0.0127133 |
| YBR162W-A | YSY6 | unknown | 0.4353 | 1.4 | 0.0094113 |
| YDR208W | MSS4 | 1-phosphatidylinositol-4-phosphate 5-kinase | 0.4397 | 1.4 | 0.286168 |
| YCR082W | AHC2 | unknown | 0.4468 | 1.4 | 0.0212216 |
| YMR002W | | unknown | 0.4538 | 1.4 | 0.0464385 |
| YPL271W | ATP15 | hydrogen-transporting ATP synthase | 0.4665 | 1.4 | 0.0300279 |
| YBR267W | REI1 | unknown | 0.4710 | 1.4 | 0.0084363 |
| YDL097C | RPN6 | structural molecule | 0.4710 | 1.4 | 0.218294 |
| YBR297W | MAL33 | transcription factor | 0.4782 | 1.4 | 0.40255 |
| YJL144W | | unknown | 0.4787 | 1.4 | 0.0291092 |
| YFL021C-A | | | 0.4793 | 1.4 | 0.222885 |
| YAL063C | FLO9 | mannose binding | 0.4820 | 1.4 | 0.0511794 |
| YLR330W | CHS5 | unknown | 0.4863 | 1.4 | 0.0320483 |
| YBR134W | | | 0.4897 | 1.4 | 0.0093899 |
| YIR016W | | unknown | 0.4918 | 1.4 | 0.134715 |
| YBR256C | RIB5 | riboflavin synthase | 0.4933 | 1.4 | 0.0786972 |
| YJL118W | | unknown | 0.5017 | 1.4 | 0.0488577 |
| YBR105C | VID24 | unknown | 0.5028 | 1.4 | 0.0489352 |
| YDL083C | RPS16B | structural constituent of ribosome | 0.5122 | 1.4 | 0.0499676 |
| YBL101W-A | | | 0.5127 | 1.4 | 0.0761509 |
| YDL005C | MED2 | RNA polymerase II transcription mediator | 0.5163 | 1.4 | 0.005962 |
| YDL212W | SHR3 | unfolded protein binding | 0.5238 | 1.4 | 0.0448283 |
| YFR008W | FAR7 | unknown | 0.5318 | 1.4 | 0.0289027 |
| YIL152W | | unknown | 0.5338 | 1.4 | 0.0120596 |
| YNL118C | DCP2 | mRNA binding | 0.5342 | 1.4 | 0.0170435 |
| YMR124W | | unknown | 0.5412 | 1.5 | 0.0749135 |
| YMR070W | MOT3 | DNA binding | 0.5435 | 1.5 | 0.0172109 |
| YDR476C | | unknown | 0.5437 | 1.5 | 0.033795 |
| YKL065C | YET1 | unknown | 0.5555 | 1.5 | 0.0510764 |
| YBR024W | SCO2 | thioredoxin peroxidase | 0.5707 | 1.5 | 0.046428 |
| YAL061W | | oxidoreductase | 0.5815 | 1.5 | 0.181938 |
| YHR198C | FMP22 | unknown | 0.5830 | 1.5 | 0.0365646 |
| YJL117W | PHO86 | unknown | 0.5858 | 1.5 | 0.0242786 |
| YDR365W-A | | | 0.5933 | 1.5 | 0.0270192 |
| YPR065W | ROX1 | specific transcriptional repressor | 0.5937 | 1.5 | 0.240102 |
| YIL063C | YRB2 | unknown | 0.5987 | 1.5 | 0.0532592 |
| YNL113W | RPC19 | DNA-directed RNA polymerase | 0.6105 | 1.5 | 0.0450949 |
| YHR154W | RTT107 | unknown | 0.6182 | 1.5 | 0.0122228 |
| YJR008W | | unknown | 0.6260 | 1.5 | 0.0528628 |
| YDR034C-C | | | 0.6343 | 1.6 | 0.0446272 |
| YCL011C | GBP2 | RNA binding | 0.6418 | 1.6 | 0.0001894 |
| YNL054W-A | | | 0.6478 | 1.6 | 0.0547295 |
| YKL172W | EBP2 | unknown | 0.6508 | 1.6 | 0.0394215 |
| YDR261W-A | | | 0.6510 | 1.6 | 0.0256109 |
| YLR410W-A | | | 0.6545 | 1.6 | 0.0468589 |
| YGL037C | PNC1 | nicotinamidase | 0.6548 | 1.6 | 0.0403934 |
| YGL237C | HAP2 | transcriptional activator | 0.6697 | 1.6 | 0.0248453 |
| YGR086C | PIL1 | protein kinase inhibitor | 0.6742 | 1.6 | 0.0354637 |
| YHR095W | | | 0.6878 | 1.6 | 0.0106695 |
| YDL020C | RPN4 | transcriptional activator | 0.7053 | 1.6 | 0.046169 |
| YJL142C | | | 0.7128 | 1.6 | 0.0471679 |
| YAR015W | ADE1 | phosphoribosylaminoimidazolesuccinocarboxamide synthase | 0.7187 | 1.6 | 0.0198603 |
| YBR034C | HMT1 | protein-arginine N-methyltransferase | 0.7282 | 1.7 | 0.0218337 |
| YDL134C-A | | | 0.7485 | 1.7 | 0.104724 |
| YGR069W | | | 0.7593 | 1.7 | 0.0024741 |
| YIL006W | YIA6 | transporter | 0.7617 | 1.7 | 0.204415 |

Table 4: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YHR138C | | endopeptidase inhibitor | 0.7718 | 1.7 | 0.0429167 |
| YCR080W | | | 0.7753 | 1.7 | 0.0319661 |
| YNL112W | DBP2 | RNA helicase | 0.7760 | 1.7 | 0.0158107 |
| YBR068C | BAP2 | amino acid transporter | 0.7792 | 1.7 | 0.0936009 |
| YOR298W | MUM3 | acyltransferase | 0.7823 | 1.7 | 0.0088567 |
| YHR166C | CDC23 | protein binding | 0.8003 | 1.7 | 0.0753333 |
| YCL067C | HMLALPHA | transcription corepressor | 0.8032 | 1.7 | 0.020451 |
| YNL143C | | | 0.8048 | 1.7 | 0.0127135 |
| YER079W | | unknown | 0.8088 | 1.8 | 0.032764 |
| YNL133C | FYV6 | unknown | 0.8183 | 1.8 | 0.0064089 |
| YKR071C | DRE2 | unknown | 0.8277 | 1.8 | 0.0887988 |
| YAL039C | CYC3 | holocytochrome-c synthase | 0.8285 | 1.8 | 0.0075338 |
| YMR121C | RPL15B | structural constituent of ribosome | 0.8292 | 1.8 | 0.0128724 |
| YEL071W | DLD3 | D-lactate dehydrogenase (cytochrome) | 0.8443 | 1.8 | 0.045225 |
| YOL143C | RIB4 | 6,7-dimethyl-8-ribityllumazine synthase | 0.8468 | 1.8 | 0.0470601 |
| YMR090W | | unknown | 0.8622 | 1.8 | 0.0460975 |
| YKR040C | | | 0.8778 | 1.8 | 0.0263266 |
| YDR203W | | | 0.8782 | 1.8 | 0.102064 |
| YBR122C | MRPL36 | structural constituent of ribosome | 0.8928 | 1.9 | 0.0345816 |
| YJR047C | ANB1 | translation initiation factor | 0.9015 | 1.9 | 0.406705 |
| YBR249C | ARO4 | 3-deoxy-7-phosphoheptulonate synthase | 0.9033 | 1.9 | 0.0395766 |
| YGR034W | RPL26B | structural constituent of ribosome | 0.9062 | 1.9 | 0.0161455 |
| YAL060W | BDH1 | (R,R)-butanediol dehydrogenase | 0.9090 | 1.9 | 0.0296579 |
| YHR187W | IKI1 | RNA polymerase II transcription elongation factor | 0.9115 | 1.9 | 0.0309065 |
| YDL048C | STP4 | unknown | 0.9160 | 1.9 | 0.0454473 |
| YDR529C | QCR7 | ubiquinol-cytochrome-c reductase | 0.9263 | 1.9 | 0.0298725 |
| YFR017C | | unknown | 0.9315 | 1.9 | 0.0450418 |
| YHL015W | RPS20 | structural constituent of ribosome | 0.9440 | 1.9 | 0.0257441 |
| YKL141W | SDH3 | succinate dehydrogenase (ubiquinone) | 0.9558 | 1.9 | 0.233962 |
| YIL144W | TID3 | structural constituent of cytoskeleton | 0.9593 | 1.9 | 0.054765 |
| YFL-TYB | | | 0.9787 | 2.0 | 0.0448005 |
| YCL065W | | | 0.9825 | 2.0 | 0.0175075 |
| YNL157W | IGO1 | unknown | 0.9865 | 2.0 | 0.0441041 |
| YKR066C | CCP1 | cytochrome-c peroxidase | 0.9895 | 2.0 | 0.0091674 |
| YNL178W | RPS3 | structural constituent of ribosome | 1.0010 | 2.0 | 0.0379982 |
| YOR020C | HSP10 | unfolded protein binding | 1.0115 | 2.0 | 0.030164 |
| YCR096C | HMRA2 | unknown | 1.0132 | 2.0 | 0.0324899 |
| YDL051W | LHP1 | RNA binding | 1.0195 | 2.0 | 0.0536717 |
| YPL058C | PDR12 | xenobiotic-transporting ATPase | 1.0252 | 2.0 | 0.20091 |
| YHR097C | | unknown | 1.0355 | 2.0 | 0.0346253 |
| YOR052C | | unknown | 1.0378 | 2.1 | 0.144021 |
| YGR183C | QCR9 | ubiquinol-cytochrome-c reductase | 1.0418 | 2.1 | 0.0751807 |
| YER032W | FIR1 | unknown | 1.0543 | 2.1 | 0.0523615 |
| YHR137W | ARO9 | aromatic-amino-acid transaminase | 1.0658 | 2.1 | 0.0942575 |
| YDL169C | UGX2 | unknown | 1.0682 | 2.1 | 0.0261166 |
| YKR049C | FMP46 | oxidoreductase | 1.0758 | 2.1 | 0.174838 |
| YMR251W-A | HOR7 | unknown | 1.0838 | 2.1 | 0.0922112 |
| YOL106W | | | 1.0888 | 2.1 | 0.0328209 |
| YNL180C | RHO5 | GTPase | 1.1487 | 2.2 | 0.028897 |
| YBR101C | FES1 | adenyl-nucleotide exchange factor | 1.1603 | 2.2 | 0.0205651 |
| YLR309C | IMH1 | unknown | 1.1613 | 2.2 | 0.0186924 |
| YDR368W | YPR1 | oxidoreductase | 1.1630 | 2.2 | 0.0361984 |
| YKL093W | MBR1 | unknown | 1.2308 | 2.3 | 0.0486059 |
| YGL062W | PYC1 | pyruvate carboxylase | 1.2363 | 2.4 | 0.0459513 |
| YBL015W | ACH1 | acetyl-CoA hydrolase | 1.2407 | 2.4 | 0.0388953 |
| YHR214C-B | | | 1.2627 | 2.4 | 0.0532289 |
| YCL040W | GLK1 | glucokinase | 1.2673 | 2.4 | 0.0381918 |
| YMR173W-A | | | 1.2678 | 2.4 | 0.0466505 |
| YHR155W | YSP1 | unknown | 1.2897 | 2.4 | 0.128931 |
| YFR016C | | unknown | 1.3243 | 2.5 | 0.0459536 |
| YHR054C | | unknown | 1.3395 | 2.5 | 0.0265463 |
| YEL009C | GCN4 | DNA binding | 1.3403 | 2.5 | 0.0382242 |
| YAR009C | | | 1.3413 | 2.5 | 0.0114855 |
| YEL012W | UBC8 | ubiquitin conjugating enzyme | 1.3488 | 2.5 | 0.0521475 |
| YDR210C-D | | | 1.3710 | 2.6 | 0.0296608 |
| YOR031W | CRS5 | copper ion binding | 1.3805 | 2.6 | 0.0265957 |
| YHR145C | | | 1.3915 | 2.6 | 0.0788772 |
| YLR256W | HAP1 | specific RNA polymerase II transcription factor | 1.3978 | 2.6 | 0.113426 |
| YHR149C | SKG6 | unknown | 1.4077 | 2.7 | 0.0385228 |

Table 4: continued.

| ORF ID | Gene Name | Gene Function | Avg Log ₂ Ratio | Fold Change | T-test |
|-----------|-----------|---|----------------------------|-------------|-----------|
| YHR070W | TRM5 | tRNA (guanine) methyltransferase | 1.4293 | 2.7 | 0.0320838 |
| TDR261C-D | | | 1.4402 | 2.7 | 0.04881 |
| YHR087W | | unknown | 1.4628 | 2.8 | 0.0438919 |
| YDR244W | PEX5 | peroxisome targeting sequence binding | 1.4683 | 2.8 | 0.0511298 |
| YER062C | HOR2 | glycerol-1-phosphatase | 1.4755 | 2.8 | 0.0441064 |
| YLR035C-A | | | 1.4912 | 2.8 | 0.0209809 |
| YGR088W | CTT1 | catalase | 1.5060 | 2.8 | 0.135822 |
| Q0060 | AI3 | endonuclease | 1.5138 | 2.9 | 0.0431973 |
| YJL116C | NCA3 | unknown | 1.5418 | 2.9 | 0.0909263 |
| YMR096W | SNZ1 | protein binding | 1.5512 | 2.9 | 0.0658355 |
| YOL053C-A | | | 1.5812 | 3.0 | 0.0462657 |
| YIL060W | | unknown | 1.5883 | 3.0 | 0.0331232 |
| YBR230C | OM14 | unknown | 1.6172 | 3.1 | 0.0502337 |
| YHR165C | PRP8 | RNA splicing factor | 1.6178 | 3.1 | 0.0352883 |
| YER024W | YAT2 | carnitine O-acetyltransferase | 1.6467 | 3.1 | 0.0457763 |
| YHR171W | ATG7 | APG12 activating enzyme | 1.6627 | 3.2 | 0.0530414 |
| YBR116C | | | 1.6698 | 3.2 | 0.0604677 |
| YHR103W | SBE22 | unknown | 1.6850 | 3.2 | 0.0194832 |
| YHR161C | YAP1801 | clathrin binding | 1.6862 | 3.2 | 0.037702 |
| YHR144C | DCD1 | dCMP deaminase | 1.7098 | 3.3 | 0.0438885 |
| YDR041W | RSM10 | structural constituent of ribosome | 1.7302 | 3.3 | 0.0532902 |
| YFL029C | CAK1 | cyclin-dependent protein kinase activating kinase | 1.7960 | 3.5 | 0.0370991 |
| YLR410W-B | | | 1.8005 | 3.5 | 0.0058466 |
| YDL182W | LYS20 | homocitrate synthase | 1.8177 | 3.5 | 0.0255799 |
| YNL110C | NOP15 | unknown | 1.8192 | 3.5 | 0.0420959 |
| YIL099W | SGA1 | glucan 1,4-alpha-glucosidase | 1.8200 | 3.5 | 0.0408364 |
| YIL053W | RHR2 | glycerol-1-phosphatase | 1.8332 | 3.6 | 0.0308083 |
| YFL021W | GAT1 | specific RNA polymerase II transcription factor | 1.9022 | 3.7 | 0.048557 |
| YOL150C | | | 1.9847 | 4.0 | 0.0398114 |
| YGR243W | FMP43 | unknown | 1.9980 | 4.0 | 0.0476662 |
| YMR174C | PAI3 | endopeptidase inhibitor | 2.0182 | 4.1 | 0.0382856 |
| YDR070C | FMP16 | unknown | 2.0265 | 4.1 | 0.0371171 |
| YGR182C | | | 2.0555 | 4.2 | 0.0106683 |
| YOL151W | GRE2 | oxidoreductase | 2.1938 | 4.6 | 0.0417326 |
| YHR139C | SPS100 | unknown | 2.1995 | 4.6 | 0.042501 |
| YCR005C | CIT2 | citrate (Si)-synthase | 2.2268 | 4.7 | 0.0099217 |
| YJL106W | IME2 | protein kinase | 2.3285 | 5.0 | 0.0508438 |
| YBR050C | REG2 | protein phosphatase type 1 regulator | 2.3793 | 5.2 | 0.0456598 |
| YLR346C | | unknown | 2.4018 | 5.3 | 0.0338526 |
| YPL135W | ISU1 | protein binding | 2.4313 | 5.4 | 0.0459425 |
| YMR195W | ICY1 | unknown | 2.4578 | 5.5 | 0.0283315 |
| YHR033W | | unknown | 2.4952 | 5.6 | 0.0476181 |
| YPL223C | GRE1 | unknown | 2.5318 | 5.8 | 0.0477036 |
| YHR162W | | unknown | 2.5717 | 5.9 | 0.0417935 |
| YBR281C | | unknown | 2.5802 | 6.0 | 0.0436952 |
| YHR056C | RSC30 | DNA binding | 2.5825 | 6.0 | 0.0005777 |
| YFL013W-A | | | 2.5992 | 6.1 | 0.0431277 |
| YHR055C | CUP1-2 | copper ion binding | 2.6320 | 6.2 | 0.0505493 |
| YFL014W | HSP12 | unknown | 2.6625 | 6.3 | 0.0461229 |
| YHR053C | CUP1-1 | copper ion binding | 2.7475 | 6.7 | 0.0360849 |
| YKL096W | CWP1 | structural constituent of cell wall | 3.1493 | 8.9 | 0.0515745 |
| YER065C | ICL1 | isocitrate lyase | 3.2103 | 9.3 | 0.0446256 |
| YMR107W | SPG4 | unknown | 3.2662 | 9.6 | 0.0388049 |
| YJR095W | SFC1 | succinate:fumarate antiporter | 3.5115 | 11.4 | 0.0203382 |
| YNL036W | NCE103 | carbonate dehydratase | 3.7867 | 13.8 | 0.0378788 |
| YCR010C | ADY2 | ammonium transporter | 3.8223 | 14.1 | 0.0387718 |
| YDR536W | STL1 | transporter | 3.9488 | 15.4 | 0.0341034 |
| YMR175W | SIP18 | phospholipid binding | 4.6685 | 25.4 | 0.0458001 |

APPENDIX II SUCCINATE METABOLISM

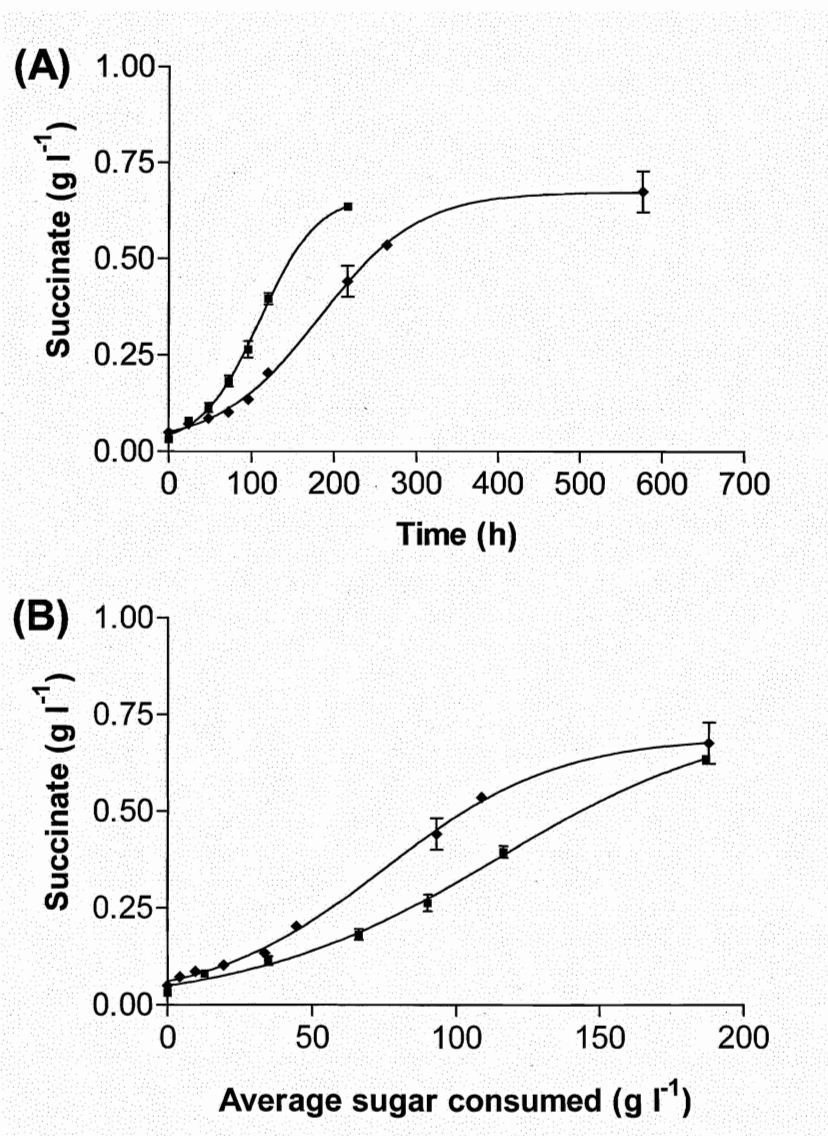


Figure 1: Yeast succinate production. The concentration of extracellular succinate (A) was measured in duplicate throughout the course of the fermentations of Icewine juice (\blacklozenge) and diluted Icewine juice (\blacksquare) and was plotted as a function of sugar consumed (B). The values represent the average \pm standard deviation of the mean of triplicate fermentations.